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(21) International Application Number: PCT/US00/11543 (22) International Filing Date: 28 April 2000 (28.04.00) (30) Priority Data: <table border="0" style="width: 100%;"><tr><td style="width: 30%;">60/131,720</td><td style="width: 40%;">30 April 1999 (30.04.99)</td><td style="width: 30%;">US</td></tr><tr><td>60/149,738</td><td>21 August 1999 (21.08.99)</td><td>US</td></tr><tr><td>60/155,945</td><td>24 September 1999 (24.09.99)</td><td>US</td></tr><tr><td>60/182,012</td><td>11 February 2000 (11.02.00)</td><td>US</td></tr></table> (71) Applicant (for all designated States except US): BIOSTRATUM, INC. [US/US]; Suite 200, 4825 Creekstone Drive, Durham, NC 27703 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): KORTESMAA, Jarrko [SE/SE]; Karolinska Institute, S-17 177 Stockholm (SE). TRYGGVASON, Karl [SE/SE]; Karolinska Institute, S-17 177 Stockholm (SE). (74) Agent: HARPER, David, S.; McDonnell, Boehnen, Hulbert & Berghoff, Suite 3200, 300 South Wacker Drive, Chicago, IL 60606 (US).		60/131,720	30 April 1999 (30.04.99)	US	60/149,738	21 August 1999 (21.08.99)	US	60/155,945	24 September 1999 (24.09.99)	US	60/182,012	11 February 2000 (11.02.00)	US	(81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>Without international search report and to be republished upon receipt of that report.</i>
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(54) Title: LAMININ 8 AND METHODS FOR ITS USE (57) Abstract The present invention provides substantially purified laminin 8, methods for making recombinant laminin 8, cells that express recombinant laminin 8, and methods for using the recombinant laminin 8 to accelerate the healing of injuries to vascular tissue and tissue of mesenchymal origin, and to promote cell attachment and migration.														

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LAMININ 8 AND METHODS FOR ITS USE

Cross Reference

This application claims priority to U.S. Provisional Patent Application Serial
5 Nos. 60/131,720 filed April 30, 1999; 60/149,738 filed August 19, 1999; 60/155,945
filed September 24, 1999; and 60/182,012 filed February 11, 2000; all of which are
incorporated herein by reference in their entirety.

Field of the Invention

10 This application relates to purified laminin 8 and methods for its use.

Background of the Invention

Basal laminae (basement membranes) are sheet-like, cell-associated
extracellular matrices that play a central role in cell growth, tissue development, and
15 tissue maintenance. They are present in virtually all tissues, and appear in the earliest
stages of embryonic development.

Basal laminae are central to a variety of architectural and cell-interactive
functions: (See for example, Malinda and Kleinman, *Int. J. Biochem. Cell Biol.* 28:957-
959 (1996); Aumailley and Krieg, *J. Invest. Dermatology* 106:209-214 (1996))

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1. They serve as architectural supports for tissues, providing adhesive substrata for cells.
2. They create perm-selective barriers between tissue compartments that impede the migration of cells and passively regulate the exchange of macromolecules.
- 25 These properties are illustrated by the kidney glomerular basement membrane, which functions as an important filtration structure, creating an effective blood-tissue barrier that is not permeable to most proteins and cells.
3. Basal laminae create highly interactive surfaces that can promote cell migration and cell elongation during embryogenesis and wound repair. Following an
30 injury, they provide a surface upon which cells regenerate to restore normal tissue function.
4. Basal laminae present information encoded in their structure to contacting cells that is important for differentiation and tissue maintenance. This information is

communicated to the cells through various receptors that include the integrins, dystroglycan, and cell surface proteoglycans. Signaling is dependent not only on the presence of matrix ligands and corresponding receptors that interact with sufficient affinities, but also on such topographical factors as ligand density in a three-dimensional matrix "landscape", and on the ability of basal lamina components to cluster receptors. Because these matrix proteins can be long-lived, basal laminae create a "surface memory" in the basal lamina for resident and transient cells.

The basal lamina is largely composed of laminin and type IV collagen heterotrimers that in turn become organized into complex polymeric structures. To date, six type IV collagen chains and at least twelve laminin subunits have been identified. These chains possess shared and unique functions and are expressed with specific temporal (developmental) and spatial (tissue-site specific) patterns.

Laminins are a family of heterotrimeric glycoproteins that reside primarily in the basal lamina. They function via binding interactions with neighboring cell receptors, and by forming laminin networks, and they are important signaling molecules that can strongly influence cellular function. Laminins are important in both maintaining cell/tissue phenotype as well as promoting cell growth and differentiation in tissue repair and development.

Laminins are large, multi-domain proteins, with a common structural organization. The laminin molecule integrates various matrix and cell interactive functions into one molecule.

The laminin molecule is comprised of an α -, β -, and γ -chain subunit joined together through a coiled-coil domain. Within this structure are identifiable domains that possess binding activity towards other laminin and basal lamina molecules, and membrane-bound receptors. Domains VI, IVb, and IVa form globular structures, and domains V, IIIb, and IIIa (which contain cysteine-rich EGF-like elements) form rod-like structures. (Kamiguchi et al., Ann. Rev. Neurosci. 21:97-125 (1998)) Domains I and II of the three chains participate in the formation of a triple-stranded coiled-coil structure (the long arm).

Table 1 shows the individual chains that each laminin type is composed of:

TABLE 1. Known laminin family members

<i>Protein</i>	<i>Chains</i>
Laminin-1	$\alpha 1\beta 1\gamma 1$
Laminin-2	$\alpha 2\beta 1\gamma 1$
Laminin-3	$\alpha 1\beta 2\gamma 1$
Laminin-4	$\alpha 2\beta 2\gamma 1$
Laminin-5	$\alpha 3\beta 3\gamma 2$
Laminin-6	$\alpha 3\beta 1\gamma 1$
Laminin-7	$\alpha 3\beta 2\gamma 1$
Laminin-8	$\alpha 4\beta 1\gamma 1$
Laminin-9	$\alpha 4\beta 2\gamma 1$
Laminin-10	$\alpha 5\beta 1\gamma 1$
Laminin-11	$\alpha 5\beta 2\gamma 1$
Laminin-12	$\alpha 2\beta 1\gamma 3$

5 Four structurally-defined family groups of laminins have been identified. The first group of five identified laminin molecules all share the $\beta 1$ and $\gamma 1$ chains, and vary by their α -chain composition ($\alpha 1$ to $\alpha 5$ chain). The second group of five identified laminin molecules all share the $\beta 2$ and $\gamma 1$ chain, and again vary by their α -chain composition. The third group of identified laminin molecules has one identified member, laminin 5, with a chain composition of $\alpha 3\beta 3\gamma 2$. The fourth group of identified laminin molecules has one identified member, laminin 12, with the newly identified $\gamma 3$ chain ($\alpha 2\beta 1\gamma 3$)

Some progress has been made in elucidating the relationship between domain structure and function. (See, for example, Wewer and Engvall, Neuromusc. Disord. 6:409-418 (1996).) The overall sequence similarity among the homologous domains in different chains varies, but it is highest in domain VI (thought to play a key role in laminin polymerization), followed by domains V (possibly involved in protein-protein interactions) and III (entactin/nidogen binding; possible cell adhesion sites), and is lowest in domains I, II (both thought to be involved in intermolecular assembly, and containing possible cell adhesion sites), and G. Not all domains are present in all 3 types of chains. The globular G domain (thought to be involved in cell receptor binding) is present only in the α chains. Other domains may not be present in all chains within a certain chain type. For example, domain VI is absent from $\alpha 3$, $\alpha 4$, and $\gamma 2$ chains. (Wewer and Engvall, 1996)

As a result of their large size (>600 kD) and unique structure, the laminin molecules can be resolved in the electron microscope. (Wewer and Engvall, 1996) Typically, laminins appear as cross-shaped molecules in an EM. The three short arms of the cross represent the amino terminal portions of each of the three separate laminin chains (one short arm per chain). The long arm of the cross is composed of the C-terminal parts of the three chains, which together form a coiled coil structure. (Wewer and Engvall, 1996) The long arm ends with the globular G domain.

The coiled-coil domain of the long arm is crucial for assembly of the three chains of laminin. (Yurchenco et al., Proc. Natl. Acad. Sci. 94:10189-10194 (1997)). Disulfide bonds bridge and stabilize all three chains in the most proximal region of the long arm and join the β and γ chains in the most distal region of the long arm.

A model of laminin receptor-facilitated self-assembly, based on studies conducted with cultured skeletal myotubes and Schwann cells, predicts that laminins bind to their receptors, which freely diffuse in a fluidic membrane, when ligand-free. Receptor engagement forces the laminins into a high local two-dimensional concentration, facilitating their mass-action driven assembly into ordered surface polymers. In this process, the engaged receptors are also reorganized, accompanied by cytoskeletal rearrangements. (Colognato, J. Cell Biol. 145:619-631 (1999)) This reorganization activates the receptors, causing signal transduction with the alteration of cell expression, shape and/or behavior. The evidence is that laminins must possess both cell-interacting and architecture-forming sites, which are located in different protein domains and on different subunits.

One class of laminin receptors are the integrins, which are cell surface receptors that mediate many cell-matrix and cell-cell interactions. Integrins are heterodimers, consisting of an α and a β subunit. 16 α - and 8 β -subunits are known, and at least 22 combinations of α and β subunits have been identified to date. Some integrins have only one or a few known ligands, whereas others appear to be very promiscuous. Binding to integrins is generally of low affinity, and is dependent on divalent cations. Integrins, activated through binding to their ligands, transduce signals via kinase activation cascades, such as focal adhesion and mitogen-activated kinases. Several different integrins bind different laminin isoforms more or less

specifically. (Aumailley et al., In The Laminins, Timpl and Ekblom, eds., Harwood Academic Publishers, Amsterdam. pp. 127-158 (1996))

Laminin 8, a recently identified laminin, is composed of $\alpha 4$, $\beta 1$, and $\gamma 1$ laminin chains. The laminin $\alpha 4$ chain is widely distributed both in adults and during
5 development. (Iivanainen et al., J. Biol. Chem. 272:27862-27868 (1997)) In adults it is found in the basement membrane surrounding cardiac, skeletal, and smooth muscle fibers, and in lung alveolar septa. Furthermore, it is found in the endothelial basement membrane both in capillaries and larger vessels, and in the perineurial basement membrane of peripheral nerves, as well as in intersinusoidal spaces, large
10 arteries, and smaller arterioles of bone marrow. (Frieser et al., Eur. J. Biochemistry 246:727-735 (1997); Miner et al., J. Cell Biol. 137:685-701 (1997); Geberhiwot et al., Exptl. Cell Res. 253:723-732 (1999); Gu et al., Blood 93:2533-2542 (1999); Iivanainen et al., J. Biol. Chem. 272:27862-27868 (1997))

Laminin 8 is a major laminin isoform in the vascular endothelium (Iivanainen
15 et al., J. Biol. Chem. 272:27862-27868 (1997); Frieser et al., 1997), is expressed and adhered to by platelets (Geberhiwot et al., Exptl. Cell Res. 253:723-732 (1999)), and is the only laminin isoform synthesized in 3T3-L1 adipocytes, with its level of synthesis shown to increase upon adipose conversion of the cells. (Niimi et al., Matrix Biology 16:223-230 (1997)) Laminin 8 was further speculated to be the
20 laminin isoform generally expressed in mesenchymal cell lineages to induce microvessels in connective tissues. (Niimi et al., 1997).

Laminin 8 has also been identified in mouse bone marrow primary cell cultures, arteriolar walls, and intersinusoidal spaces where data indicated that it is the major laminin isoform in the developing bone marrow. (Gu et al., Blood 93:2533-
25 2542 (1999). The investigators concluded that, due to its localization in adult bone marrow adjacent to hematopoietic cells, laminin isoforms containing the $\alpha 4$ chain are the most likely to have biologically relevant interactions with developing hematopoietic cells. (Gu et al., 1999)

Despite the broad tissue distribution of the laminin $\alpha 4$ chain and laminin 8,
30 there is not a means to prepare substantially purified laminin 8 from cell or tissue sources for research and therapeutic purposes, nor has a means for recombinant expression of laminin 8 been developed. Such research and therapeutic purposes

include, but are not limited to, methods for treating injuries to tissue of mesenchymal origin, such as bone, cartilage, tendon, and ligament, treating injuries to vascular tissue, promoting cell attachment and migration, promoting therapeutic angiogenesis and neural regeneration, ex vivo cell therapy, improving the biocompatibility of medical devices, and preparing improved cell culture devices and media.

Thus, there is a need in the art for adequate amounts of substantially purified laminin-8 for research and therapeutic purposes, and methods for making laminin 8. Such laminin 8 could be prepared either from cell lines in culture, or via recombinant DNA technology.

A preferred method of production is the use of recombinant DNA technology to engineer a human cell line of choice to produce recombinant laminin-8 ("r-laminin 8"). A recombinant-based method of laminin-8 production has several advantages over purification from human tissue or isolation from human cell lines in culture:

1. The recombinant produced protein is free of human pathogens. While this is also true for endogenous cell culture produced protein, protein derived from human tissue carries a risk for contamination by HIV, hepatitis, and other infectious agents.

2. Expression levels of the protein, and hence yields, can be improved through the use of genetically engineered genes/vectors that enhance the production of the encoded protein.

3. It is possible to engineer additional peptide sequences to the protein chain that provides a binding site for a commercially viable affinity purification procedure.

4. The method can provide for the modification of protein structure/function through the addition, substitution, elimination, and/or other modifications of protein domain structures. For example, it may be desirable to introduce an integrin binding site (e.g. RGD), switch integrin recognition sites, or engineer in a stable binding site to a synthetic substrate. Thus, the creation of expression vectors that express laminin chains generates enormous flexibility for future uses and creates a basis for creating second generation "designer" laminins.

Summary of the Invention

The present invention fulfills the need in the art for a source of substantially purified laminin 8 protein, methods for making substantially purified recombinant laminin 8 (hereinafter referred to as r-laminin 8), pharmaceutical compositions comprising laminin 8, and methods of using laminin 8 for treating injuries to tissue of mesenchymal origin, such as bone, cartilage, tendon, and ligament, treating injuries to vascular tissue, promoting cell attachment and migration, ex vivo cell therapy, improving the biocompatibility of medical devices, and preparing improved cell culture devices and media.

10 In one aspect, the present invention provides recombinant host cells that express laminin 8 chains and secrete r-laminin 8. In another aspect, the present invention provides substantially purified laminin 8, and methods for producing substantially purified r-laminin 8.

In a further aspect, the present invention provides pharmaceutical compositions, comprising laminin 8 together with a pharmaceutically acceptable carrier. Such pharmaceutical compositions can optionally be provided with other extracellular matrix components.

In further aspect, the present invention provides methods and kits for accelerating the healing of injuries to tissue of mesenchymal origin, such as bone, cartilage, tendon, and ligament, treating injuries to vascular tissue, and for improving the biocompatibility of grafts used for treating such injuries. In specific examples, laminin 8 or pharmaceutical compositions thereof are used to:

- a. promote re-endothelialization at the site of vascular injuries;
- b. improve the "take" of grafts;
- 25 c. improve the biocompatibility of medical devices;
- d. treat neural injuries (neural regeneration);
- e. regulate angiogenesis; and
- d. promote cell attachment and migration

by providing an amount effective of r-laminin 8 for the various methods. In preferred embodiments of all of these methods, recombinant laminin 8 is used. The kits comprise an amount of laminin 8 effective for the desired effect, and instructions for the use thereof.

In a further aspect, the present invention provides improved medical devices and grafts, and methods for preparing improved medical devices and grafts, wherein the improvement comprises applying an amount effective of laminin 8 or the pharmaceutical compositions of the invention to the device or graft for the desired application.

In a further aspect, the invention provides improved cell culture devices, and methods for preparing improved cell culture devices, for the growth and maintenance of cells in culture, by providing an amount effective of laminin 8 for the attachment of cells to a cell culture device for the subsequent proliferation/differentiation/stasis of the cells.

In another aspect, the invention provides a cell culture growth supplement, comprising laminin 8. In another aspect, the invention provides an improved cell culture growth media, wherein the improvement comprises the addition of r-laminin 8.

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Brief Description of the Figures

Figure 1 is a photograph of a 3-12% gradient SDS-PAGE gel. LN-1 is laminin 1/nidogen (ndg) complex with component chain identities indicated on the left; LN-8 is recombinant laminin 8. Interpretation of r-laminin 8 protein band identities are indicated based on western blotting data: $\alpha 4$ = reactivity with anti-human laminin $\alpha 4$ and also anti-FLAG monoclonal antibody (mAb); $\beta 1$ = reactivity with polyclonal anti-murine laminin $\alpha 1/\beta 1/\gamma 1$; $\gamma 1$ = reactivity with anti-human laminin $\gamma 1$ mAb; * = reactivity with both anti-laminin $\gamma 1$ mAb and anti-murine $\alpha 1/\beta 1/\gamma 1$. Both samples were run on the same gel which was subsequently silver stained.

Figure 2 is a rotary shadowed electron micrograph of r-laminin 8. Top: low magnification field showing several r-laminin 8 molecules. Bottom: Individual molecules. Each molecule has two short arms and one long arm. In some molecules, a very short (5-10 nm) rod-like stub can be seen at the junction of the arms. Arrow: G-domain can be seen as consisting of two moieties in some molecules. (Bar = 50 nm)

Figure 3 is a graph depicting a titration of cell adhesion to r-laminin 8 or laminin 1.

Figure 4 is a graph depicting HT-1080 cell adhesion to r-laminin 8 or laminin 1 coated at 10 $\mu\text{g/ml}$ on 96 well plates in the presence and absence of various function-blocking integrin antibodies and other compounds.

Figure 5 is a graph depicting bovine capillary endothelial (BCE) cell adhesion to r-laminin 8 or laminin 1 coated at 10 $\mu\text{g/ml}$ on 96 well plates in the presence and absence of various function-blocking integrin antibodies and other compounds.

Figure 6 is a graph depicting immortal mouse brain endothelial (IBE) cell adhesion to r-laminin 8 or laminin 1 coated at 10 $\mu\text{g/ml}$ on 96 well plates in the presence and absence of various function-blocking integrin antibodies and other compounds.

Figure 7 is a graph depicting integrin $\alpha 6\beta 4$ -transfected K562 cell adhesion to r-laminin 8 or laminin 1 coated at 10 $\mu\text{g/ml}$ on 96 well plates in the presence and absence of various function-blocking integrin antibodies and other compounds.

Figure 8 is a graph depicting integrin $\alpha 6$ -transfected K562 cell adhesion to r-laminin 8 or laminin 1 coated at 10 $\mu\text{g/ml}$ on 96 well plates in the presence and absence of various function-blocking integrin antibodies and other compounds.

Detailed Description of the Preferred Embodiments

All references, patents and patent applications are hereby incorporated by reference in their entirety.

Within this application, unless otherwise stated, the techniques utilized may be found in any of several well-known references such as: *Molecular Cloning: A Laboratory Manual* (Sambrook, et al., 1989, Cold Spring Harbor Laboratory Press), *Gene Expression Technology* (Methods in Enzymology, Vol. 185, edited by D. Goeddel, 1991. Academic Press, San Diego, CA), "Guide to Protein Purification" in *Methods in Enzymology* (M.P. Deutscher, ed., (1990) Academic Press, Inc.); *PCR Protocols: A Guide to Methods and Applications* (Innis, et al. 1990. Academic Press, San Diego, CA), *Culture of Animal Cells: A Manual of Basic Technique, 2nd Ed.* (R.I. Freshney. 1987. Liss, Inc. New York, NY), *Gene Transfer and Expression Protocols*, pp. 109-128, ed. E.J. Murray, The Humana Press Inc., Clifton, N.J.), and the Ambion 1998 Catalog (Ambion, Austin, TX).

As used herein "laminin 8" encompasses both r-laminin 8 and heterotrimeric laminin 8 from naturally occurring sources.

As used herein, the term "r-laminin 8" refers to recombinant heterotrimeric laminin 8, expressed by a cell that has been transfected with one or more expression vectors comprising at least one nucleic acid sequence encoding a laminin 8 chain selected from the $\alpha 4$, $\beta 1$ and $\gamma 1$ chains, or a portion of the chains that are capable of forming a heterotrimeric laminin 8 and maintaining laminin 8 activity, or processed forms thereof. Such r-laminin 8 can thus comprise $\alpha 4$, $\beta 1$, and $\gamma 1$ sequences from a single organism, or from different organisms. Various laminin 8 chain DNA sequences are known in the art, and the use of each to prepare the r-laminin 8 of the invention is contemplated. (See, for example, Iivanainen et al., FEBS Letters 365:183-188 (1995); Frieser et al., Eur. J. Biochem. 246:727-735 (1997); Richards et al., Eur. J. Biochem. 238:813-821 (1996); Liu and Mayne, 15:433-437 (1996); Vuolteenaho et al., J. Biol. Chem. 265:15611-15616 (1990); Kallunki et al., J. Biol. Chem. 266:221-228 (1991); Sasaki et al., J. Biol. Chem. 263:16536-16544 (1988); Sasaki and Yamada, J. Biol. Chem. 262:17111-17117 (1987); Sasaki et al., Proc. Natl. Acad. Sci. 84:935-939 (1987); Pikkarainen et al., J. Biol. Chem. 262:10454-10462 (1987); all references incorporated by reference herein in their entirety).

The invention encompasses those laminin molecules wherein one or two of the chains that make up the recombinant heterotrimeric laminin 8 are encoded by endogenous laminin 8 chains. In a preferred embodiment, cells are transfected with one or more expression vectors comprising nucleic acid sequences encoding each of the $\alpha 4$, $\beta 1$ and $\gamma 1$ chains, or a portion of each of the chains that are capable of forming a heterotrimeric laminin 8 and maintaining laminin 8 activity.

In the present invention, laminin 8 is a secreted protein, which is capable of being directed to the ER, secretory vesicles, and the extracellular space as a result of a signal sequence, as well as those proteins released into the extracellular space without necessarily containing a signal sequence. If the secreted protein is released into the extracellular space, the secreted protein can undergo extracellular processing to produce a "mature" protein. Such processing event can be variable, and thus may yield different versions of the final "mature protein". The substantially purified laminin 8 of the present invention includes heterotrimers comprising both the full length and any such processed laminin 8 chains.

As used herein, the term "substantially purified" means that the laminin 8 so designated has been separated from its in vivo cellular environment.

As used herein, a laminin 8 polypeptide chain refers to a polypeptide chain according to one or more of the following:

5 (a) comprises a polypeptide structure selected from the group consisting of:

1. R1-R2-R3
2. R1-R2-R3(e)
3. R3
4. R3(e)
- 10 5. R1-R3
6. R1-R3(e)
7. R2-R3
8. R2-R3(e)

wherein R1 is an amino terminal methionine; R2 is a signal sequence
15 that is capable of directing secretion of the polypeptide, wherein the signal sequence may be the natural signal sequence for the particular laminin chain, that of another secreted protein, or an artificial sequence; R3 is a secreted laminin chain selected from the $\alpha 4$, $\beta 1$, and $\gamma 1$ chains; and R3(e) is a secreted laminin chain selected from the $\alpha 4$, $\beta 1$, and $\gamma 1$ chains that further comprises an epitope tag (such as those described below),
20 which can be placed at any position within the laminin chain amino acid sequence; and/or

(b) is encoded by a polynucleotide that is substantially similar to one or more of the disclosed laminin chain polynucleotide sequences or portions thereof (SEQ ID NOS.: 1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, or 27); and/or

25 (c) is encoded by a polynucleotide that hybridizes under high or low stringency conditions to the coding regions, or portions thereof, of one or more of the recombinant laminin 8 chain DNA sequences disclosed herein (SEQ ID NOS.: 1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, 27), or complementary sequences thereof; and/or

(d) has at least 70% identity to one or more of the disclosed laminin 8
30 polypeptide chain amino acid sequences (SEQ ID NOS.: 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, or 28), preferably at least 80% identity, and most preferably at least about 90% identity.

The phrase "substantially similar" is used herein in reference to polynucleotide or polypeptide sequences having one or more conservative variations from the laminin 8 sequences disclosed herein, including but not limited to deletions, insertions, inversions, repeats, and substitutions, wherein the resulting laminin chain is functionally equivalent to those disclosed herein.

For example, conservative polynucleotide variants may contain alterations in coding regions, non-coding regions, or both. Especially preferred are polynucleotide variants containing alterations which produce silent substitutions, additions, or deletions, but do not alter the properties or activities of the encoded polypeptide. Nucleotide variants produced by silent substitutions due to the degeneracy of the genetic code are preferred. Moreover, variants in which 5-10, 1-5, or 1-2 amino acids are substituted, deleted, or added in any combination are also preferred. Polynucleotide variants can be produced for a variety of reasons, including but not limited to optimizing codon expression for a particular host (change codons in the human mRNA to those preferred by a bacterial host such as *E. coli*).

Naturally occurring conservative variants are called "allelic variants," and refer to one of several alternate forms of a gene occupying a given locus on a chromosome of an organism. (Genes II, Lewin, B., ed., John Wiley & Sons, New York (1985).) These allelic variants can vary at either the polynucleotide and/or polypeptide level. Alternatively, non-naturally occurring conservative variants may be produced by mutagenesis techniques or by direct synthesis.

Using known methods of protein engineering and recombinant DNA technology, conservative polynucleotide variants may be generated to improve or alter the characteristics of the expressed laminin chain polypeptides. For instance, one or more amino acids can be deleted from the N-terminus or C-terminus of the secreted protein. (See, for example, Ron et al., J. Biol. Chem. 268: 2984-2988 (1993); Dobeli et al., J. Biotechnology 7:199-216 (1988)) Ample evidence demonstrates that variants often retain a biological activity similar to that of the naturally occurring protein. (See, for example, Gayle et al., J. Biol. Chem 268:22105-22111 (1993)) Furthermore, even if deleting one or more amino acids from the N-terminus or C-terminus of a polypeptide results in modification or loss of one or more biological functions, other biological activities may still be retained.

Guidance concerning how to make phenotypically silent amino acid substitutions is provided in Bowie, J. U. et al., Science 247:1306-1310 (1990), wherein

the authors indicate that there are two main strategies for studying the tolerance of an amino acid sequence to change.

The first strategy exploits the tolerance of amino acid substitutions by natural selection during the process of evolution. By comparing amino acid sequences in different species, conserved amino acids can be identified. These conserved amino acids are likely important for protein function. In contrast, the amino acid positions where substitutions have been tolerated by natural selection indicates that these positions are not critical for protein function. Thus, positions tolerating amino acid substitution could be modified while still maintaining biological activity of the protein.

The second strategy uses genetic engineering to introduce amino acid changes at specific positions of a cloned gene to identify regions critical for protein function. For example, site directed mutagenesis or alanine-scanning mutagenesis (introduction of single alanine mutations at every residue in the molecule) can be used. (Cunningham and Wells, Science 244:1081-1085 (1989).) The resulting mutant molecules can then be tested for biological activity.

As the authors state, these two strategies have revealed that proteins are surprisingly tolerant of amino acid substitutions. The authors further indicate which amino acid changes are likely to be permissive at certain amino acid positions in the protein. For example, most buried (within the tertiary structure of the protein) amino acid residues require nonpolar side chains, whereas few features of surface side chains are generally conserved. Moreover, tolerated conservative amino acid substitutions involve replacement of the aliphatic or hydrophobic amino acids Ala, Val, Leu and Ile; replacement of the hydroxyl residues Ser and Thr; replacement of the acidic residues Asp and Glu; replacement of the amide residues Asn and Gln, replacement of the basic residues Lys, Arg, and His; replacement of the aromatic residues Phe, Tyr, and Trp, and replacement of the small-sized amino acids Ala, Ser, Thr, Met, and Gly.

The "substantially similar" polypeptides of the present invention also include (i) substitutions with one or more of the non-conserved amino acid residues, where the substituted amino acid residues may or may not be one encoded by the genetic code, or (ii) substitution with one or more amino acid residues having substituents groups, or (iii) fusion of the mature polypeptide with another compound, such as a compound to increase the stability and/or solubility of the polypeptide (for example, polyethylene glycol), or (iv) fusion of the polypeptide with additional amino acids, such as an IgG Fc fusion region peptide, or leader or secretory sequence, or a sequence facilitating

purification. Such variant polypeptides are deemed to be within the scope of those skilled in the art from the teachings herein.

For example, polypeptide variants containing amino acid substitutions of charged amino acids with other charged or neutral amino acids may produce proteins with improved characteristics, such as less aggregation. Aggregation of pharmaceutical formulations both reduces activity and increases clearance due to the aggregate's immunogenic activity. (Pinckard et al., Clin. Exp. Immunol. 2:331-340 (1967); Robbins et al., Diabetes 36: 838-845 (1987); Cleland et al., Crit. Rev. Therapeutic Drug Carrier Systems 10:307-377 (1993).)

“Stringency of hybridization” is used herein to refer to conditions under which nucleic acid hybrids are stable. The invention also includes nucleic acids that hybridize under high stringency conditions (as defined herein) to all or a portion of the coding sequences of the laminin chain polynucleotides disclosed herein, or their complements. The hybridizing portion of the hybridizing nucleic acids is typically at least 50 nucleotides in length. As known to those of skill in the art, the stability of hybrids is reflected in the melting temperature (T_M) of the hybrids. T_M decreases approximately 1-1.5°C with every 1% decrease in sequence homology. In general, the stability of a hybrid is a function of sodium ion concentration and temperature. Typically, the hybridization reaction is performed under conditions of lower stringency, followed by washes of varying, but higher, stringency. Reference to hybridization stringency relates to such washing conditions. Thus, as used herein, high stringency refers to an overnight incubation at 42° C in a solution comprising 50% formamide, 5x SSC (750 mM NaCl, 75 mM sodium citrate), 50 mM sodium phosphate (pH 7.6), 5x Denhardt's solution, 10% dextran sulfate, and 20 µg/ml denatured, sheared salmon sperm DNA, followed by washing the filters in 0.1x SSC at about 65°C.

Also contemplated are laminin 8-encoding nucleic acid sequences that hybridize to the polynucleotides of the present invention at lower stringency hybridization conditions. Changes in the stringency of hybridization and signal detection are primarily accomplished through the manipulation of formamide concentration (lower percentages of formamide result in lowered stringency); salt conditions, or temperature. For example, lower stringency conditions include an overnight incubation at 37°C in a solution comprising 6X SSPE (20X SSPE = 3M NaCl; 0.2M NaH_2PO_4 ; 0.02M EDTA,

pH 7.4), 0.5% SDS, 30% formamide, 100 ug/ml salmon sperm blocking DNA; followed by washes at 50°C with 1XSSPE, 0.1% SDS. In addition, to achieve even lower stringency, washes performed following stringent hybridization can be done at higher salt concentrations (e.g. 5X SSC).

5 Note that variations in the above conditions may be accomplished through the inclusion and/or substitution of alternate blocking reagents used to suppress background in hybridization experiments. Typical blocking reagents include Denhardt's reagent, BLOTTO, heparin, denatured salmon sperm DNA, and commercially available proprietary formulations. The inclusion of specific blocking
10 reagents may require modification of the hybridization conditions described above, due to problems with compatibility.

As used herein, "percent identity" of two amino acids or of two nucleic acids is determined using the algorithm of Karlin and Altschul (Proc. Natl. Acad. Sci. USA 87:2264-2268, 1990), modified as in Karlin and Altschul (Proc. Natl. Acad. Sci. USA
15 90:5873-5877, 1993). Such an algorithm is incorporated into the NBLAST and XBLAST programs of Altschul et al. (J. Mol. Biol. 215:403-410, 1990). BLAST nucleotide searches are performed with the NBLAST program, score = 100, wordlength = 12, to obtain nucleotide sequences homologous to the nucleic acid molecules of the invention. BLAST protein searches are performed with the XBLAST program, score =
20 50, wordlength = 3, to obtain an amino acid sequence homologous to a polypeptide of the invention. To obtain gapped alignments for comparison purposes, Gapped BLAST is utilized as described in Altschul et al. (Nucleic Acids. Res. 25:3389-3402, 1997). When utilizing BLAST and Gapped BLAST programs, the default parameters of the respective programs (e.g., XBLAST and NBLAST) are used. See
25 <http://www.ncbi.nlm.nih.gov>.

Further embodiments of the present invention include polynucleotides encoding laminin 8 chain polypeptides having at least 70% identity, preferably at least 80% identity, and most preferably at least 90% identity to one or more of the polypeptide sequences contained in SEQ ID NO:2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, or
30 fragments thereof.

As used herein, "α4 polynucleotide" refers to polynucleotides encoding an α4 laminin chain of the same name. Such polynucleotides can be characterized by one or

more of the following: (a) the nucleotides of said polynucleotide may encode an amino acid sequence substantially similar to the sequence set forth in SEQ ID NO: 2, 4, 6, 8, 10, 12 or fragments thereof; (b) polynucleotides that encode polypeptides which share at least 70% identity, preferably 80% identity, and most preferably at least 90% identity with the sequence set forth in SEQ ID NO: 2, 4, 6, 8, 10, 12, or fragments thereof; (c) the $\alpha 4$ polynucleotides hybridize under low or high stringency conditions to the coding sequence set forth in one or more of SEQ ID NO: 1, 3, 5, 7, 9, 11, or fragments thereof, or complementary sequences thereof; or (d) the $\alpha 4$ polynucleotides encode a polypeptide with a general structure selected from (1) R1-R2-R3; (2) R1-R2-R3(e); (3) R3; (4) R3(e); (5) R1-R3; (6) R1-R3(e); (7) R2-R3; and (8) R2-R3(e); wherein R1 and R2 are as described above, and R3 and R3(e) are as described above but comprise secreted $\alpha 4$ chain polypeptides.

As used herein, " $\beta 1$ polynucleotides" refers to polynucleotides encoding a $\beta 1$ laminin chain of the same name. Such polynucleotides can be characterized by one or more of the following: (a) the nucleotides of said polynucleotides may encode an amino acid sequence substantially similar to the sequence set forth in SEQ ID NO: 14, 16, 18, 20 or fragments thereof; (b) polynucleotides that encode polypeptides which share at least 70% identity, preferably at least 80%, and most preferably at least 90% identity with one or more of the sequences set forth in SEQ ID NO: 14, 16, 18, 20 or fragments thereof; (c) the $\beta 1$ polynucleotides hybridize under low or high stringency conditions to the coding sequence set forth in one or more of SEQ ID NO: 13, 15, 17, 19, fragments thereof, or complementary sequences thereof; or (d) the $\beta 1$ polynucleotides encode a polypeptide with a general structure selected from (1) R1-R2-R3; (2) R1-R2-R3(e); (3) R3; (4) R3(e); (5) R1-R3; (6) R1-R3(e); (7) R2-R3; and (8) R2-R3(e); wherein R1 and R2 are as described above, and R3 and R3(e) are as described above but comprise secreted $\beta 1$ chain polypeptides.

As used herein, " $\gamma 1$ polynucleotides" refers to polynucleotides encoding a $\gamma 1$ laminin chain of the same name. Such polynucleotides can be characterized by one or more of the following: (a) the nucleotides of said polynucleotides may encode an amino acid that is substantially similar to one or more of the sequences set forth in SEQ ID NO: 22, 24, 26, 28 or fragments thereof; (b) polynucleotides that encode polypeptides which share at least 70% identity, preferably at least 80%, and most preferably at least

90% identity with at one or more of the sequences set forth in SEQ ID NO: 22, 24, 26, 28 or fragments thereof; (c) the $\gamma 1$ polynucleotides hybridize under low or high stringency conditions to the coding sequence set forth in one or more of SEQ ID NO: 21, 23, 25, 27 or complementary sequences thereof; or (d) the $\gamma 1$ polynucleotides
5 encode a polypeptide with a general structure selected from (1) R1-R2-R3; (2) R1-R2-R3(e); (3) R3; (4) R3(e); (5) R1-R3; (6) R1-R3(e); (7) R2-R3; and (8) R2-R3(e); wherein R1 and R2 are as described above, and R3 and R3(e) are as described above but comprise secreted $\gamma 1$ chain polypeptides.

As used herein, the term "epitope tag" refers to a polypeptide sequence that is
10 expressed as part of a chimeric protein, where the epitope tag serves as a recognition site for binding of antibodies generated against the epitope tag, or for binding of other molecules that can be used for affinity purification of sequences containing the tag.

As used herein, the term "increased biocompatibility" refers to reduced induction of acute or chronic inflammatory response, and reduced disruption of the
15 proper differentiation of implant-surrounding tissues for laminin 8-coated biomaterials relative to an analogous, non-coated biomaterial.

As used herein the term "graft" refers to both natural and prosthetic grafts and implants.

In one aspect, the present invention provides r-laminin 8 expressing-cells that
20 have been transfected with an expression vector containing promoter sequences that are operatively linked to nucleic acid sequences encoding at least one polypeptide sequence comprising the $\alpha 4$, $\beta 1$ and $\gamma 1$ chains of laminin 8, or fragments thereof, wherein the transfected cells secrete heterotrimeric laminin 8 containing the recombinant laminin chain. In a preferred embodiment, the cells are systematically transfected with
25 recombinant expression vectors containing promoter sequences that are operatively linked to nucleic acid sequences encoding polypeptide sequences comprising the $\alpha 4$, $\beta 1$ and $\gamma 1$ chains of laminin 8. After the multiple transfections, the cells express each of the recombinant laminin 8 chains, which form the heterotrimer, before r-laminin 8 secretion into the media.

30 In a preferred embodiment, cDNAs encoding the $\alpha 4$, $\beta 1$ and $\gamma 1$ chains, or fragments thereof, are subcloned into an expression vector. Alternatively, laminin 8 $\alpha 4$, $\beta 1$ and/or $\gamma 1$ gene sequences, including one or more introns, can be used.

Any cell capable of expressing and secreting the r-laminin 8 can be used. Preferably, eukaryotic cells are used, and most preferably mammalian cells are used, including but not limited to kidney and epithelial cell lines. In a most preferred embodiment, the mammalian cells do not express all of the laminin 8 chains endogenously. Carbohydrate and disulfide post-translational modifications are believed to be required for laminin 8 protein folding and function. This makes the use of eukaryotic cells preferable for producing functional r-laminin 8, although other systems are useful for obtaining, for example, antigens for antibody production.

"Recombinant expression vector" includes vectors that operatively link a nucleic acid coding region or gene to any promoter capable of effecting expression of the gene product. The promoter sequence used to drive expression of the individual chains or r-laminin 8 may be constitutive (driven by any of a variety of promoters, including but not limited to, CMV, SV40, RSV, actin, EF) or inducible (driven by any of a number of inducible promoters including, but not limited to, tetracycline, ecdysone, steroid-responsive). The expression vector must be replicable in the host organisms either as an episome or by integration into host chromosomal DNA. In a preferred embodiment, the expression vector comprises a plasmid. However, the invention is intended to include other expression vectors that serve equivalent functions, such as viruses.

In one embodiment, at least one of the laminin chain polypeptide sequences, or fragments thereof, is operatively linked to a nucleic acid sequence encoding an "epitope tag", so that at least one of the chains is expressed as a fusion protein with an expressed epitope tag. The epitope tag may be expressed as the amino terminus, the carboxy terminus, or internal to any of the polypeptide chains comprising r-laminin 8, so long as the resulting r-laminin 8 remains functional. Any epitope tag may be utilized, so long as it can be used as the basis for affinity purification of the resulting r-laminin 8. Examples of such epitope tags include, but are not limited to FLAG (Sigma Chemical, St. Louis, MO), myc (9E10) (Invitrogen, Carlsbad, CA), 6-His (Invitrogen; Novagen, Madison, WI), and HA (Boehringer Mannheim Biochemicals).

In another embodiment, one of the r-laminin 8 chains is expressed as a fusion protein with a first epitope tag, and at least one other r-laminin chain is expressed as a fusion protein with a second epitope tag. This permits multiple rounds of purification

to be carried out. Alternatively, the same epitope tag can be used to create fusion proteins with more than one of the r-laminin chains.

In a further embodiment, the epitope tag can be engineered to be cleavable from the r-laminin 8 chain(s). Alternatively, no epitope tag is fused to any of the r-laminin 8 chains, and the r-laminin 8 is purified by standard techniques, including but not limited to affinity chromatography using laminin 8 specific antibodies or other laminin 8 binding molecules.

Transfection of the expression vectors into eukaryotic cells can be accomplished via any technique known in the art, including but not limited to calcium phosphate co-precipitation, electroporation, or liposome mediated-, DEAE dextran mediated-, polycationic mediated-, or viral mediated transfection. Transfection of bacterial cells can be done by standard methods.

In a preferred embodiment, the cells are stably transfected. Methods for stable transfection and selection of appropriate transfected cells are known in the art. In a most preferred embodiment, a CMV promoter driven expression vector is used in a human kidney embryonic 293 cell line.

Media from cells transfected with a single laminin chain are initially analyzed on Western blots using laminin chain-specific antibodies. The expression of single laminin chains following transfection is generally intracellular. Clones showing reactivity against individual transfected chain(s) are verified by any appropriate method, such as PCR, reverse transcription-PCR, or nucleic acid hybridization, to confirm incorporation of the transfected gene. Preferably, analysis of genomic DNA preparations from such clones is done by PCR using laminin chain-specific primer pairs. Media from transfected clones producing all three chains are further analyzed for r-laminin 8 secretion and/or activity, by any appropriate method, including Western blot analysis and cell binding assays. Activity of the r-laminin 8 is preferably analyzed in a cell adhesion assay.

In another aspect, the present invention provides substantially purified laminin 8, preferably r-laminin 8. In one embodiment, the substantially purified laminin 8 comprises a first chain comprising an $\alpha 4$ chain polypeptide; a second chain comprising a $\beta 1$ chain polypeptide; and a third chain comprising a $\gamma 1$ chain polypeptide. Alternatively, the r-laminin 8 comprises a first chain that is substantially similar to at

least one of the sequences shown in SEQ ID NO: 2, 4, 6, 8, 10, 12 or fragments thereof; a second chain that is substantially similar to at least one of the sequence shown in SEQ ID NO: 14, 16, 18, 20 or fragments thereof; and a third chain that is substantially similar to the sequence shown in SEQ ID NO: 22, 24, 26, 28 or fragments thereof.

5 In another embodiment, the substantially purified r-laminin 8 comprises a first chain comprising a polypeptide that is at least about 70% identical to at least one of the sequences shown in SEQ ID NO: 2, 4, 6, 8, 10, 12 or fragments thereof; a second chain comprising a polypeptide that is at least 70% identical to at least one of the sequences shown in SEQ ID NO: 14, 16, 18, 20 or fragments thereof; and a third chain comprising
10 a polypeptide that is at least 70% identical to at least one of the sequences shown in SEQ ID NO: 22, 24, 26, 28 or fragments thereof, wherein the first, second, and third polypeptides are produced recombinantly, and wherein the first, second, and third chains assemble into a recombinant heterotrimeric laminin 8.

In a preferred embodiment, at least one of the first, second, or third chains of the
15 substantially purified human r-laminin 8 is expressed as a fusion protein with an epitope tag.

Alternatively, the r-laminin 8 comprises a heterotrimeric polypeptide structure, wherein each individual chain comprises a general structure selected from the group consisting of: (1) R1-R2-R3; (2) R1-R2-R3(e); (3) R3; (4) R3(e); (5) R1-R3; (6) R1-
20 R3(e); (7) R2-R3; and (8) R2-R3(e)

wherein R1 is a amino terminal methionine; R2 is a signal sequence that is capable of directing secretion of the polypeptide, wherein the signal sequence may be the natural signal sequence for the particular laminin chain, that of another secreted protein, or an artificial sequence; R3 is a secreted $\alpha 4$, $\beta 1$, or $\gamma 1$ laminin chain; and
25 R3(e) is a secreted laminin $\alpha 4$, $\beta 1$, and $\gamma 1$ chain that further comprises an epitope tag (such as those described above), which can be placed at any position within the laminin chain amino acid sequence.

In a preferred embodiment, purification of r-laminin 8 is accomplished by passing media from the transfected cells through an antibody affinity column. In one
30 embodiment, antibodies against a peptide epitope expressed on at least one of the recombinant chains are attached to an affinity column, and bind the r-laminin 8 that has been secreted into the media. The r-laminin 8 is removed from the column by passing

excess peptide over the column. Eluted fractions are analyzed by any appropriate method, including gel electrophoresis and Western blot analysis. In a further embodiment, the peptide epitope can be cleaved after purification. In other embodiments, two or three separate r-laminin chains are expressed as fusion proteins, each with a different epitope tag, permitting two or three rounds of purification and a doubly or triply purified r-laminin 8. The epitope tag can be engineered so as to be cleavable from the r-laminin 8 chain(s) after purification. Alternatively, no epitope tag is fused to any of the r-laminin 8 chains, and the r-laminin 8 is purified by standard techniques, including but not limited to affinity chromatography using laminin 8 specific antibodies or other laminin 8 binding molecules.

The present invention further provides pharmaceutical compositions comprising substantially purified laminin 8 and a pharmaceutically acceptable carrier. In a preferred embodiment, the pharmaceutical composition comprises substantially purified r-laminin 8. According to this aspect of the invention, other agents can be included in the pharmaceutical compositions, depending on the condition being treated. The pharmaceutical composition may further comprise one or more other compounds, including but not limited to any of the collagens, other laminin types, fibronectin, vitronectin, cadherins, integrins, α -dystroglycan, entactin/nidogen, α -dystroglycan, glycoproteins, proteoglycans, heparan sulfate proteoglycan, glycosaminoglycans, epidermal growth factor, vascular endothelial growth factor, fibroblast growth factor, or nerve growth factors, and peptide fragments thereof.

Pharmaceutical preparations comprising substantially purified laminin 8 can be prepared in any suitable form, and generally comprise the laminin 8 in combination with any of the well known pharmaceutically acceptable carriers. The carriers can be injectable carriers, topical carriers, transdermal carriers, and the like. The preparation may advantageously be in a form for topical administration, such as an ointment, gel, cream, spray, dispersion, suspension or paste. The preparations may further advantageously include preservatives, antibacterials, antifungals, antioxidants, osmotic agents, and similar materials in composition and quantity as is conventional. Suitable solutions for use in accordance with the invention are sterile, are not harmful for the proposed application, and may be subjected to conventional pharmaceutical operations such as sterilization and/or may contain conventional adjuvants, such as preservatives,

stabilizers, wetting agents, emulsifiers, buffers etc. For assistance in formulating the compositions of the present invention, one may refer to Remington's Pharmaceutical Sciences, 15th Ed., Mack Publishing Co., Easton, Pa. (1975).

In further aspect, the present invention provides methods and kits comprising
5 laminin 8, or pharmaceutical compositions thereof (and instructions for using the laminin 8 in the kits) for accelerating the healing of injuries to tissue of mesenchymal origin, such as bone, cartilage, tendon, and ligament, treating injuries to vascular and neural tissue, and for improving the biocompatibility of grafts used for treating such injuries. In a preferred embodiment of each of the methods disclosed below, r-laminin
10 8 is used. In specific examples, substantially purified laminin 8, r-laminin 8, or pharmaceutical compositions thereof are used to:

- a. promote re-endothelialization at the site of vascular injuries;
- b. improve the "take" of grafts;
- c. improve the biocompatibility of medical devices;
- 15 d. treat neural injuries (neural regeneration);
- e. regulate angiogenesis; and
- d. promote cell attachment and migration

by providing an amount effective of laminin 8 or pharmaceutical compositions thereof for the various methods.

20 In one embodiment, laminin 8 is used to promote re-endothelialization, and to thus inhibit abnormal smooth muscle cell proliferation, at the site of a vascular injury. The $\alpha 4$ chain is associated with mesenchymally derived cell populations, including but not limited to endothelium and smooth muscle cells, and laminin 8 has been shown to be a primary laminin of the vascular endothelium.

25 The value of angioplasty in clearing occluded coronary arteries is limited by a restenosis/reocclusion rate of 50-70%. Several studies have indicated that the insertion of a vascular stent following angioplasty appears to decrease the occurrence of restenosis, but the problem still limits the effectiveness of this treatment. Restenosis appears to arise in part from the proliferation of vascular smooth muscle
30 cells in response to the angioplasty treatment. It is likely that the scraping action of angioplasty removes not only the problematic occlusion, but also sections of the vascular basal lamina. The discontinuous basal lamina that results could contribute to

what appears to be abnormal growth of the vascular smooth muscle cells that leads to restenosis.

The attachment of laminin 8 to vascular stents can be used to limit restenosis, by promoting re-endothelialization. The interaction of vascular endothelial cells with the laminin 8 coated stents promotes their adhesion and attachment, thereby leading to homeostasis and a normal cell growth response, instead of the injury/activation endothelial cell response seen with restenosis. While activated platelets adhere to laminin 8, non-activated platelets do not. Furthermore, it has been shown that soluble laminin 8 does not cause platelet activation, but has an inhibitory effect on platelet activation by classical activators such as thrombin, collagen I, and ADP (unpublished observations). A more normal and controlled rate of re-endothelialization will decrease the incidence of re-occlusions, and improve the outcome of the angioplasty procedure.

Similarly, synthetic vascular grafts can induce blood clotting and thrombosis through interactions of blood clotting factors with the synthetic graft material. Coating vascular grafts with laminin 8 promotes endothelialization of the synthetic vessel, thereby providing for a non-thrombogenic surface. Vascular endothelial cells, like other cells that sit upon a basement membrane, prefer to adhere to an appropriate basement membrane substrate. Laminin-8 has been identified as a component of the vascular basal lamina, and is suspected to be involved in the attachment of vascular endothelial cells to the supporting basal lamina. Providing this substrate in a graft material creates a non-thrombogenic surface, promotes endothelialization, and inhibits intravascular thrombosis and vascular obstruction.

Administration to the injured blood vessel can be accomplished in some cases by simply coating laminin 8 or pharmaceutical compositions thereof into an injured area. In other embodiments, delivery can be accomplished by:

1. Coating a stent;
2. Coating a biodegradable sleeve over the stent; or
3. Forcing a liquid preparation of the laminin 8 or pharmaceutical compositions thereof through a porous catheter to the injured site.

In another embodiment, the present invention provides methods to promote bone and connective tissue repair in a subject. The incorporation of laminin 8 or pharmaceutical compositions thereof into wound repair dressings and matrices as well

as tissue grafts to accelerate the healing of bone and connective tissue repair provides a natural ligand interactive surface to promote normal cell adherence, cell growth and tissue development. Many grafts are used to replace connective tissue that has a cell layer adherent to a basal lamina. When an inappropriate surface is provided to these
5 cells following grafting, the graft is at risk for failure of restoration of the normal cell layer. The advantage of coating these grafts with laminin 8 is to create a surface that sufficiently recapitulates a normal basal lamina surface to promote cell re-population. As used herein the term "graft" refers to both natural and prosthetic grafts.

The methods of the present invention have application in the healing of tendon,
10 cartilage, or ligament tears, deformities and defects, bone fractures, defects, as well as use in the improved fixation of tendon, cartilage, or ligament to bone or other tissues. In addition, bony in-growth into various prosthetic devices can be greatly enhanced so that such artificial parts are firmly and permanently anchored into the surrounding skeletal tissue through a natural osseous bridge.

15 In a further aspect, the present invention comprises medical devices with improved biocompatibility, wherein the devices are coated with laminin 8 or pharmaceutical compositions thereof, alone or in combination with other proteins or agents that serve to increase the biocompatibility of the device surface. The coated device stimulates cell attachment and provides for diminished inflammation and/or
20 infection at the site of entry of the appliance.

Such medical devices can be of any material used for implantation into the body, and preferably are made of or coated with a biocompatible metal that may be either stainless steel or titanium. Alternatively, the device is made of or coated with a ceramic material, or a polymer including but not limited to polyester, polyglycolic acid
25 or a polygalactose-polyglycolic acid copolymer.

One particular use of the present invention is to increase cell adhesion to target surfaces, including but not limited to endothelial, skeletal muscle, smooth muscle, and other mesenchymally-derived cells. For example, vascular grafts and stents may be coated with laminin 8 or pharmaceutical compositions thereof to stimulate endothelial
30 cell attachment. Alternatively, bone or connective tissue grafts or prostheses may be coated with laminin 8 or pharmaceutical compositions thereof to stimulate adhesion of the appropriate cell type and improved grafting efficiency.

If the device is made of a natural or synthetic biodegradable material in the form of a mesh, sheet or fabric, laminin 8 or pharmaceutical compositions thereof may be applied directly to the surface thereof. Appropriate cells may then be cultured on the matrix to form transplantable or implantable devices, including dental abutment pieces, needles, metal pins or rods, indwelling catheters, colostomy tubes, surgical meshes and any other appliance for which coating with laminin 8 is desirable. Alternatively, the devices may be implanted and cells may be permitted to attach in vivo.

Coupling of the substantially purified laminin 8 may be non-covalent (such as by adsorption), or by covalent means. The device may be immersed in, incubated in, or sprayed with the laminin 8 or pharmaceutical compositions thereof.

The dosage regimen for various treatments using the laminin 8 of the present invention is based on a variety of factors, including the type of injury or condition, the age, weight, sex, medical condition of the individual, the severity of the condition, and the route of administration. Thus, the dosage regimen may vary widely, but can be determined routinely by a physician using standard methods. Laminins are extremely potent molecules, and one or a few molecules per cell could produce an effect. Thus, effective doses in the pico-gram per milliliter range are possible if the delivery is optimized. Laminins are sometimes present in an insoluble form in the basement membrane and have the capability of polymerizing at concentrations as low as about 50 $\mu\text{g/ml}$, depending on the laminin isoform and the conditions. Laminins can also polymerize into a gel at a concentration of about 2-3 mg/ml . Dosage levels of the order of between 1 ng/ml and 10 mg/ml are thus useful for all methods disclosed herein, preferably between about 1 $\mu\text{g/ml}$ and about 3 mg/ml .

The present invention also provides a method for inducing cell attachment to the device (as disclosed above), comprising coating the appliance with laminin 8 or pharmaceutical compositions thereof prior to incubation with cells appropriate for the desired application.

Laminin preparations are known to induce the growth and differentiation of neurons (U.S. Patent No. 5,229,365), and have been used in combination with Type I collagen to coat a hollow conduit and promote nerve regeneration across a gap of severed nerve. (U.S. Patent No. 5,019,087)

Thus, in another embodiment, a method is provided for nerve regeneration, comprising administering to a subject in need thereof an amount effective of laminin 8 or pharmaceutical compositions thereof to promote nerve regeneration. The graft can comprise a nerve graft, or a prosthetic graft. Both bioresorbable and non-resorbable materials have been used in tubes for bridging nerve gaps. (See for example, Nyilas, et al., (Trans. Soc. Biomater., 6, 85, 1983), Molander, et al. (Biomaterials, Vol. 4, pp. 276-280, October, 1983), Colin, et al., (Journal of Dental Research July, 1984, pp. 987-993). The method can be used to treat diseases and injuries characterized by the loss of function and or/degeneration of neurons and nerves.

Laminins, or cell extracts containing laminins have been shown to regulate angiogenesis in a biphasic manner. (See, for example, Nicosia et al., Dev. Biol. 164:197-206 (1994); Bonfil et al., Int. J. Cancer 58:233-239 (1994)). At lower concentrations (30-300 $\mu\text{g/ml}$), a laminin-entactin complex stimulated angiogenesis in a three-dimensional culture, while at 3000 $\mu\text{g/ml}$ the same complex was inhibitory to angiogenesis. Thus, in another aspect, the present invention provides methods for regulating angiogenesis, comprising contacting a tissue or culture substrate with an amount effective of laminin 8 or pharmaceutical compositions thereof to regulate angiogenesis. In one embodiment, the laminin 8 is used to promote angiogenesis by contacting a tissue or culture substrate with an amount effective of laminin 8 to promote angiogenesis. In another embodiment, the laminin 8 is used to inhibit angiogenesis, by contacting the tissue or culture substrate with an amount effective of laminin 8 to inhibit angiogenesis. An example of culture substrates to be contacted with laminin 8 to regulate angiogenesis are those used for tissue engineering purposes.

In another aspect of the present invention, laminin 8 is used for the culture of cells, including but not limited to endothelial cells, nerve cells, cells of hematopoietic lineage, and mesenchymally-derived cells including but not limited to cells derived from bone, connective tissue, and adipose tissue, skeletal muscle cells, and smooth muscle cells, by contacting the cells with an amount effective of laminin 8 to stimulate attachment and proliferation/differentiation/stasis of cells. The laminin 8 can either be provided in the cell culture medium, or as a cell culture medium supplement, or may be coated on the surface of a cell growth substrate. In a preferred embodiment, the method further includes contacting the cells with other compounds, including but not

limited to any of the collagens, other laminin types, fibronectin, α -dystroglycan, cadherins, integrins, entactin/nidogen, α -dystroglycan, glycoproteins, proteoglycans, heparan sulfate proteoglycan, glycosaminoglycans, epidermal growth factor or nerve growth factors, vascular endothelial growth factor, fibroblast growth factor, and peptide
5 fragments thereof.

The cells may comprise primary cells or cell culture cell lines. The methods of this aspect of the invention can be used in vivo, ex vivo, or in vitro.

In a preferred embodiment, laminin 8 is used to coat the surface of a substrate, to promote cell adhesion to the substrate, and to stimulate cell
10 proliferation/differentiation/stasis. The substrate used herein may be any desired substrate. For laboratory use, the substrate may be as simple as glass or plastic. For use in vivo, the substrate may be any biologically compatible material capable of supporting cell adhesion. Suitable substrate materials include shaped articles made of or coated with such materials as collagen, regenerated collagen, polyglycolic acid,
15 polygalactose, polylactic acid or derivatives thereof; biocompatible metals such as titanium and stainless steel; ceramic materials including prosthetic material such as hydroxylapatite; synthetic polymers including polyesters and nylons; polystyrene; polyacrylates; polytetrafluoroethylene and virtually any other material to which biological molecules can readily adhere. The determination of the ability of a particular
20 material to support adhesion of the r-laminin 8 of the invention requires only routine experimentation by the skilled artisan.

In a further aspect, the present invention provides cell growth substrates for adhesion and culturing of cells, by providing an amount effective of laminin 8 for the attachment of cells to a cell culture device for the attachment and subsequent
25 proliferation/differentiation/stasis of the cells. The substrates may comprise any of the substrates discussed above.

In another aspect of the present invention, an improved cell culture medium is provided, wherein the improvement comprises addition to the cell culture medium of an effective amount of laminin 8 to the cell culture medium to promote the adherence,
30 proliferation, and/or maintenance of cells. Any cell culture media that can support the growth of cells can be used with the present invention. Such cell culture media include, but are not limited to Basal Media Eagle, Dulbecco's Modified Eagle Medium, Iscove's

Modified Dulbecco's Medium, McCoy's Medium, Minimum Essential Medium, F-10 Nutrient Mixtures, Opti-MEM® Reduced-Serum Medium, RPMI Medium, and Macrophage-SFM Medium or combinations thereof.

5 The improved cell culture medium can be supplied in either a concentrated (ie: 10X) or non-concentrated form, and may be supplied as either a liquid, a powder, or a lyophilizate. The cell culture may be either chemically defined, or may contain a serum supplement. Culture media is commercially available from many sources, such as GIBCO BRL (Gaithersburg, MD) and Sigma (St. Louis, MO). In an alternative embodiment, the laminin 8 is used as a cell culture supplement.

10 The laminin 8 or pharmaceutical compositions thereof of the present invention can be used for the treatment of a variety of conditions and diseases as described herein, including but not limited to various vascular, neural, and mesenchymal tissue injuries, including but not limited to angioplasty restenosis, tissue ischemia, neural damage, vascular surgical procedures, atherosclerosis, bone fractures, defects, and
15 disorders which result in weakened bones such as osteoporosis, osteoarthritis, and periodontal disease; bone loss resulting from cancer or side effects of other medical treatment; age-related loss of bone mass; articular cartilage tears, deformities and other cartilage defects such as arthritis and cartilaginous tissue damage, tendon or ligament tears, deformities and other tendon or ligament defects such as tendinitis and carpal
20 tunnel syndrome, periodontal ligament injury, and tendon-to-bone detachment.

The amount of laminin 8 or pharmaceutical compositions thereof used in such treatments will, of course, depend upon the type and severity of the condition or disease being treated, the route of administration chosen, and will be determined by the attending physician or veterinarian. The term "therapeutically effective amount" of
25 laminin 8 or pharmaceutical compositions thereof refers to the amount of laminin 8 or pharmaceutical compositions thereof, in the absence of other exogenously applied factors, determined to produce a therapeutic response in a mammal. Such therapeutically effective amounts are readily ascertained by one of ordinary skill in the art.

30 The present invention may be better understood with reference to the accompanying examples that are intended for purposes of illustration only and should

not be construed to limit the scope of the invention, as defined by the claims appended hereto.

EXAMPLES

5 *Expression Constructs*

For expression of the human laminin $\alpha 4$ chain containing a C-terminal FLAG epitope, the full length cDNA was constructed and modified as follows. Complementary DNA lambda clones subcloned into pBluescriptTM or pCRscriptTM (Stratagene) plasmid vectors from an earlier study (Iivanainen et al., 1995) were used
10 as cDNA source, except for clone FL136. The EcoRI insert from FL136 lambda DNA was cloned into the pBluescriptTM EcoRI site to make FL136E. The 0.78 kb SacI-BamHI fragment from clone FL76 was ligated into SacI-BamHI digested pSL1180 (Pharmacia) to make FL76SB. A sequence corresponding to nucleotides 2378-4274 of human laminin $\alpha 4$ cDNA was PCR-amplified using cDNA library as a template,
15 digested with SacI and cloned into the FL76SB SacI site and its orientation confirmed to make HL4-SB. The FL64 BamHI-Sall fragment was cloned into HL4-SB BamHI-Sall to make HL4-3'.

The Eco72I-XhoI fragment from clone FL117 was ligated into the Eco72I-XhoI sites of FL136E to make HL4-5'. Both mouse and human laminin $\alpha 4$ cDNAs have
20 poorly conserved Kozak-sequences at the translation initiation site, as well as several extra 5' untranslated region (UTR) ATG sequences. To ensure efficient and correct translation initiation, the Kozak sequence was edited to match the consensus and the rest of the 5' UTR was deleted using standard molecular biology techniques. The resulting product was EcoRI-EagI-digested and cloned to the EcoRI-EagI-digested
25 HL4-5' to make HL4Mut-5'. The SpeI-XhoI fragment from HL4Mut-5' was cloned into HL4-3' to make clone HL4-Full with full length cDNA. The EcoRI insert from HL4-Full was cloned into pcDNA3.1/Zeo(-) expression vector (Invitrogen) to make HL4-Full.pcDNA. (SEQ ID NO:1)

The sequence encoding the FLAG epitope (SEQ ID NO:3) was inserted as
30 follows. The FL64 BamHI-HindIII fragment was cloned into pUC19 to make FL64BH. PCR was performed using primers to introduce the FLAG epitope, using HL4-3' as template. The product was digested with XbaI and HindIII and cloned into

XbaI-HindIII digested FL64BH to make HL4FLAG-3'. This also resulted in deletion of the original 3' UTR. The BamHI-HindIII fragment from HL4FLAG-3' was cloned into BamHI-HindIII-digested HL4-Full.pcDNA vector, replacing the original BamHI-HindIII fragment to make HL4FLAG-B, which lacked the BamHI-BamHI fragment.

5 The final expression construct named HL4FLAG-Full was made by inserting the missing BamHI fragment in the correct orientation. All PCR-derived parts of the cDNA sequence were sequenced to ensure that no mutations had occurred during amplification.

The construct used for expression of the mouse laminin β 1 chain (SEQ ID
10 NO:15) has been previously described (Yurchenco et al., *Proc. Natl. Acad. Sci. U. S. A.* 94(19), 10189-94 (1997)).

To make the construct named HG1 for expression of the human laminin γ 1 chain, full length cDNA (SEQ ID NO:19) encoding the human laminin γ 1 chain was released with BamHI from a baculovirus expression vector pVL941 (unpublished) and
15 cloned into the BamHI site of a pcDNA3.1/Hygro(-) mammalian expression vector (Invitrogen).

Antibodies, control proteins, and cell lines

Affinity purified polyclonal anti-laminin α 4 antibody (Ab) S8 was prepared as
20 described previously. (Iivanainen et al., 1997, *J. Biol. Chem.* 272(44), 27862-8) Polyclonal anti-EHS-laminin Ab, anti-FLAG M2 monoclonal Ab (mAb), purified control mouse IgG, RGDS-peptide and heparin (grade I-A) were purchased from Sigma Chemical Company (St. Louis, MO). Anti-laminin γ 1 (clone 22) mAb was from Transduction Laboratories (Lexington, KY). Mouse function blocking mAbs against
25 integrin α 1 (clone FB12), integrin α 2 (clone P1E6), and integrin α 3 (clone P1B5) were obtained from Chemicon (Temecula, CA). Rat function blocking mAbs anti-integrin α 6 (clone GoH3) and control rat IgG_{2a} were also from Chemicon. Rat function blocking mAbs against integrin α 5 (clone BIIG2) and integrin β 1 (clone AIIB2) were provided by Dr. C. Damsky (Univ. of California, San Francisco) as hybridoma
30 supernatants. Immunoglobulins were purified from the supernatants using GAMMABIND PLUSTM Sepharose (Pharmacia; Stockholm, Sweden) according to the manufacturer's instructions. Secondary Ab conjugates anti-rabbit IgG-HRP and anti-

mouse IgG-HRP were from Dakopatts (Denmark). Laminin 1 from EHS-tumor, collagen type IV from EHS-tumor, and human placental laminin were obtained from Sigma. Fibronectin and some of the laminin 1 from EHS-tumor were purchased from Gibco BRL (Rockville, MD). EHS-derived laminin 1/nidogen complex was kindly
5 provided by Dr. J. Engel (Univ. of Basel, Switzerland). Human fibrosarcoma HT-1080 (CCL-121) cells were from the American Type Tissue Collection. (Manassas, VA) IMMORTOMOUSETM brain capillary endothelial (IBE, Kanda et al., 1999, *Exp. Cell Res.* 248(1), 203-13) and bovine adrenal microvascular (BCE, Folkman et al., 1979) cells were kindly provided by Dr. L. Claesson-Welsh (Medical Biochemistry and
10 Microbiology, Univ. of Uppsala) and K. Olausson (Medical Cell Biology, Univ. of Uppsala). Three human erythroleukemic K562 cell lines transfected to express integrins $\alpha 3$ (Delwel et al., 1994, *Mol. Biol. Cell* 5(2), 203-15), $\alpha 6$ (Delwel et al., 1993, *J. Biol. Chem.* 268(34), 25865-75), or both $\alpha 6$ and $\alpha 4$ (Niessen et al., 1994, *Exp. Cell Res.* 211(2), 360-7 *Mol. Biol. Cell*) were provided by Dr. A. Sonnenberg (Netherlands
15 Cancer Institute, Amsterdam, Netherlands).

Production and purification of recombinant laminin 8

Recombinant laminin 8 ("r-laminin 8") was produced in human embryonic kidney cells (HEK-293, ATCC CRL-1573) cultured in DME/pyruvate/10% fetal calf
20 serum (FCS) at 37°C in a humidified 5% CO₂ atmosphere. Wild-type cells were stably transfected with the laminin $\beta 1$ expression construct as previously described (Yurchenco et al., 1997) and selected using 500 μ g/ml G418. All further cell culture and clonal expansion was carried out in the continuous presence of relevant selection antibiotics. A highly expressing clone was then transfected with the HL4FLAG-Full
25 construct using standard calcium-phosphate transfection methods, and stable colonies were selected using 300 μ g/ml Zeocin. Clones were isolated using cloning rings, expanded, and analyzed for laminin $\alpha 4$ secretion by Western blotting of medium using the anti-laminin $\alpha 4$ Ab S8. The clone with the highest expression was transfected with the HG1 construct, and stable clones were selected using 100 μ g/ml hygromycin.
30 These clones were then screened via Western blotting using a mAb against laminin $\gamma 1$, and clones showing the highest secretion were expanded further.

For production of r-laminin 8, cells were grown in the culture medium for up to four days, after which the medium was collected and centrifuged to remove cell debris. After collection, Tris-Cl pH 7.5 was added to 50 mM and EDTA was added to a concentration of 10 mM. If not used immediately, the medium was stored at -70°C.

- 5 For protein production into serum-free medium, confluent cultures were washed twice with PBS and the medium was changed to DME supplemented with pyruvate, insulin-transferrin-selen supplement (Sigma) and 1 µg/ml aprotinin (Sigma).

r-laminin 8 was affinity purified using an anti-FLAG M2 matrix (Sigma). Before use, the matrix was washed with 0.1M glycine (pH 3.5) and TBS (50 mM Tris-
10 HCl pH 7.5/150 mM NaCl) according to the manufacturer's instructions. Brij-20 (Fluka, Milwaukee, Wisconsin) was added to the medium to a final concentration of 0.05% (v/v), and the medium was incubated in batch mode with the matrix (25 µl matrix/ml) overnight at 4°C with agitation. The matrix was collected by passing the medium through a sintered column, and washed extensively in the column first with
15 TBS/1 mM EDTA and then PBS/1 mM EDTA. Bound r-laminin 8 was competitively eluted with 100 µg/ml FLAG peptide (Sigma) in PBS/1mM EDTA at room temperature. The matrix was then regenerated as recommended by the manufacturer. The eluate was diluted 1:1 with 20 mM NaPO₄/1 mM EDTA (pH 7.5), and injected into a UNO-Q ion-exchange column (Bio-Rad, Hercules, CA). At this salt concentration,
20 the FLAG peptide passes through, but r-laminin 8 is bound. The column was then washed with 20 mM phosphate/1 mM EDTA, and the r-laminin 8 was eluted with 20 mM NaPO₄/1.5M NaCl/1 mM EDTA. The eluate was diluted 1:10 with 20 mM NaPO₄ (pH 7.5)/1 mM EDTA to a final salt concentration of 150 mM and concentrated using 100 kD cut-off ultrafiltration (Gelman; Ann Arbor, Michigan) to approximately 0.5
25 mg/ml.

Characterization of r-laminin 8

Secreted r-laminin 8 in cell medium and after purification was characterized using linear 5% or 6% SDS-PAGE and 3-12% gradient SDS-PAGE under reducing and
30 non-reducing conditions. Proteins were visualized using silver staining or blotted to PVDF membranes using a semi-dry blotting system (Bio-Rad). For immunodetection, Renaissance ECL-System (Dupont, Waltham, MA) was used in conjunction with the

Abs described above. Protein quantitation was done by measuring absorbance at 280 nm or using the Bradford method (Bio-Rad protein assay kit).

Rotary shadowing electron micrography (EM) was performed as described previously. (Yurchenco and Chen, 1993) When purifying for rotary shadowing, the
5 matrix was equilibrated with 0.15 M NH_4HCO_3 -acetate buffer (pH 7.4), and the r-laminin 8 was then eluted with FLAG-peptide in the same buffer.

Adhesion assays and cell culture

For adhesion assays, flat-bottom 96 well plates (Maxi-Sorp, Nunc; Rochester,
10 NY) were coated by incubating with proteins diluted in PBS overnight at 4°C (50 μl /well). The remaining protein-binding capacity was saturated by addition of 2% heat-inactivated BSA in PBS (50 μl /well) and further incubation for at least 4 hours. Prior to assaying, the coating/blocking solution was aspirated, and the wells were washed with the binding medium (100 μl /well). Drying of coated protein was avoided, since
15 this was found to be detrimental to adhesion in some cases.

All cells were cultured in humidified 5% CO_2 atmosphere. HT-1080 and BCE cells were cultured in DME/10% FCS/pyruvate at 37°C, BCE on gelatin-coated plastic. IBE cells were cultured in F12/10% FCS/2 U/ml γ -interferon on gelatin-coated plastic at 33°C. Transfected K562 cells were grown in suspension in RPMI/10%FCS
20 supplemented with 1 mg/ml G418 at 37°C. For K562 cells transfected with both $\alpha 4$ and $\beta 4$ integrins, 0.7 mg/ml hygromycin was included in the medium. Prior to dissociation, the cells were washed twice with PBS. HT-1080 cells were disassociated using 5 mM EDTA in PBS, while the others were disassociated using trypsin-EDTA (Gibco-BRL). To remove trypsin, cells were pelleted and resuspended twice in serum-
25 free medium. Cells were counted and suspended in buffered serum-free medium at $2-3 \times 10^5$ cells/ml. K562 cells were washed twice with serum-free medium and resuspended at 10^6 cells/ml. DME/25 mM HEPES/pyruvate was used for HT-1080 cells; F12/25 mM HEPES/0.25% BSA was used for other cell types.

K562 cell stimulation was done using 5 ng/ml PMA (Sigma). Antibodies or
30 other test compounds were added to the cell suspension and the cells were allowed to recover at 37°C for 30 minutes. The cells were then added to the protein-coated 96-

well plates (100 μ l/well) and allowed to adhere for 30 (K562) or 60 (other cells) minutes at 37°C. To remove unbound cells, wells were washed by two or three cycles of careful addition of 100 μ l of binding medium followed by aspiration. The remaining cells were fixed with 1% glutaraldehyde in PBS for 10 minutes at room temperature.

5 Cells were stained with 0.1% crystal violet (Sigma) for 30 minutes and unbound stain was removed by four washes with water. Bound stain was solubilized in 2% SDS (100 μ l/well) and quantitated by measuring the absorbance at 595 nm using a microplate reader.

None of the cell lines bound appreciably to BSA. When the quantitative results were calculated, binding to BSA was given a value of zero, while the relevant control was given the value of 100. The mean and SEM were calculated from results obtained from parallel wells.

RESULTS

15 *Production and characterization of r-laminin 8*

Unconcentrated medium from wild-type HEK-293 cells did not react in Western blots with the anti-laminin α 4, anti-laminin γ 1, anti-EHS-laminin, or anti-FLAG antibodies, indicating that these cells express endogenous laminins at very low amounts if at all. The transfected α 4 chain could be secreted to some extent even when expressed alone, but secretion of the other chains required simultaneous expression of all three chains. Cells transfected with laminin α 4, β 1, and γ 1 chain expression constructs secreted large amounts of all three chains to the medium. The best cell clones ("G1-2" and "G1-3") were estimated to produce 3-5 milligrams of r-laminin 8 per liter of medium.

25 The r-laminin 8 bound to anti-FLAG M2 matrix with high specificity. When eluted competitively with the FLAG peptide, only laminin α 4, β 1, and γ 1 bands were seen in silver-stained 3-12% gradient SDS-PAGE gels. (Figure 1) Under non-reducing conditions, the purified protein hardly entered the gel, which was to be expected as the predicted molecular weight for the mature trimer is at least 570 kD. A minor fraction of the purified trimer appeared as non-covalently associated (see discussion). In this fraction, the β 1 and γ 1 chains appeared as covalently associated dimers, whereas the α 4 chain was non-covalently associated. Under reducing conditions, the protein appeared

30

as a broad band at around 200 kD, which reacted on Western blots with $\alpha 4$, EHS, $\gamma 1$, and anti-FLAG antibodies. The predicted molecular weights for mature $\alpha 4$, $\beta 1$, and $\gamma 1$ polypeptides are 200, 195, and 174 kD respectively. Laminins are heavily glycosylated, which may account for the slight discrepancy in molecular weight observed in SDS-PAGE. The $\beta 1$ and $\gamma 1$ chains of laminin 1 purified from EHS-tumor showed similar or slightly slower mobility than those of r-laminin 8.

Rotary shadowing EM revealed r-laminin 8 to be a Y-shaped molecule with two short and one long arm in accordance with the predicted structure. (Figure 2) In many cases, a very short (5-10 nm) rod-like stub could be seen at the junction of the arms. The G-domains could sometimes be seen as consisting of two moieties.

Cell binding to r-laminin 8 and receptor identification

We assayed the binding of human fibrosarcoma (HT-1080) and transfected K562 cells to r-laminin 8 in the presence of different blocking anti-integrin antibodies to identify integrin receptors binding to r-laminin 8. Immortomouse brain capillary endothelial (IBE) and bovine adrenal microvascular endothelial (BCE) cells were also used to study the adhesion of endothelial cells to r-laminin 8. The BCE cells express at least integrins $\alpha 6\beta 1$, $\alpha 6\beta 4$, $\alpha 1\beta 1$, $\alpha 2\beta 1$, $\alpha 3\beta 1$, $\alpha 5\beta 1$, $\alpha v\beta 1$, $\alpha v\beta 3$, and $\alpha 5\beta 5$ (Klein et al., 1993, *Mol. Biol. Cell* 4(10), 973-82), whereas IBE cells have been reported to express integrins $\alpha 3$, $\alpha 5$ and $\beta 1$, but not $\alpha 1$, $\alpha 2$, or $\alpha 6$ (Kanda et al., 1999)

When compared to laminin 1, the adhesiveness of r-laminin 8 was quantitatively similar or slightly weaker for the cell lines studied, as approximately the same number of cells bound to both substrates after washing. (Figure 3) Similar results were obtained with IBE and BCE cells (data not shown). In further cell adhesion assays (Figures 4-8), cells were allowed to bind to either-laminin 8 or laminin 1 coated at 10 $\mu\text{g/ml}$ on 96 well plates. Prior to the assay, different components were added to the cell medium. Values indicated are relative to that of control antibody (normal mouse IgG for mouse antibodies and rat IgG_{2b} for rat monoclonal antibodies), which was designated as 100. For other substances, the same volume of buffer was added. Adhesion to bovine serum albumin (BSA) was designated zero. The text under the columns indicate the integrin subunit blocked or the added substance. Error bars

indicate SEM. Integrin monoclonal antibodies were used at 10 μ g/ml, heparin at 2 mg/ml, and EDTA at 5 mM.

Monoclonal antibodies against integrins α 1 and α 2 were tested only in the HT-1080 cell line, where they had no or only small effects on cell binding, indicating that these integrins were not major mediators of adhesion to r-laminin 8 (Figure 4). Adhesion to Type IV collagen was reduced to about 50% by the anti-integrin α 2 mAb, demonstrating the presence of active α 2 integrin (data not shown). Integrin α 1 mAb had only a slight effect on adhesion to collagen IV when used alone, but it had a synergistic effect when used in combination with the α 2 mAb (not shown). The Ab against integrin α 3 had only minor effects on adhesion of HT-1080 when used alone, but it had a synergistic effect when used in combination with the α 2 mAb (data not shown). The monoclonal antibody to α 2 integrin had only minor effects on adhesion of HT 1080 (Figure 4) cells to fibronectin or laminin, even though the cells have been shown to express high levels of α 3 β 1 (Wayner et al., 1993, *J. Cell Biol.* 121(5), 1141-52). The blocking of integrin α 5 had a slight stimulating effect on HT-1080 adhesion to both laminin 1 and laminin 8 (Figure 4), whereas adhesion of BCE cells to r-laminin 8 was slightly reduced (Figure 5). The mAb did block adhesion of HT-1080 cells to fibronectin almost completely, indicating the presence of active α 5 integrin in these cells (not shown).

α -6 subunit containing integrin(s) were identified as the major mediators of adhesion to r-laminin 8. The integrin α 6 subunit is known to associate with either β 1 or β 4 (Sonnenberg et al., 1990, *J. Cell Sci.* 96(Pt 2), 207-17). By using a mAb (GoH3) that blocks α 6 β 1- and α 6 β 4-mediated binding, we could completely abolish binding of HT-1080 and BCE cells to r-laminin 8. An anti- β 1 integrin mAb (AIIIB2) also completely blocked the binding of HT-1080 cells to r-laminin 8 indicating that integrin α 6 β 1 is crucial for adhesion of these cells to r-laminin 8 (Figure 4). In contrast, binding of BCE cells was blocked only partially (about 70%) by the anti- β 1 mAb, suggesting that these endothelial cells use both α 6 β 1 and α 6 β 4 to adhere to r-laminin 8 (Figure 5). In another endothelial cell line, the mouse IBE cells, the anti- α 6 subunit mAb blocked the binding to r-laminin 8 only partially (about 60%), suggesting that the

cells are using, in addition to $\alpha 6$ -subunit containing integrins, also other receptors (Figure 6).

Interestingly, when adhesion to r-laminin 8 was compared to that of laminin 1, it was observed that the adhesion was quite differently affected by the blocking anti- $\alpha 6$ and anti- $\alpha 1$ integrin mAbs. HT-1080 cells interacted with laminin 1 not only via $\alpha 6\beta 1$ integrin, but also via other $\beta 1$ -subunit-containing integrin(s), since the blocking was only partial with anti- $\alpha 6$, but complete with anti- $\beta 1$. (Figure 4) Furthermore, the adhesion of BCE cells to laminin 1 was mediated by $\beta 1$ integrin(s) other than $\alpha 6\beta 1$, since the adhesion was completely blocked by anti- $\beta 1$, but was only minimally affected by anti- $\alpha 6$. (Figure 5) Similarly, in IBE cells, the adhesion to laminin 1 was mediated by receptors other than $\alpha 6$ integrin(s), since it was not affected by anti- $\alpha 6$. (Figure 6)

To verify the role of $\alpha 6\beta 1$ and $\alpha 6\beta 4$ integrins as r-laminin 8 receptors, transfected K562 cells were used. Parental K562 cells endogenously express only integrin $\alpha 5\beta 1$, which is in an inactive state. The cells normally grow in suspension but can be made adherent with an activating anti- $\beta 1$ Ab or stimulation with PMA. K562 cells transfected with the $\alpha 6$ subunit express $\alpha 6\beta 1$ on the cell surface (Delwel et al., 1993). Interestingly, while these cells bound laminin 1 efficiently only after stimulation with PMA, they bound r-laminin 8 strongly even without stimulation (Figure 8). This finding demonstrates that the adhesive properties of r-laminin 8 are different from those of laminin 1. The cell adhesion to both laminin isoforms could be blocked with either anti-integrin $\alpha 6$ or $\beta 1$ mAbs, which agrees with results obtained with other cell lines (Figure 4-5). In addition to inactive $\alpha 6\beta 1$, K562 cells transfected with $\alpha 6$ and $\beta 4$ subunits express constitutively active $\alpha 6\beta 4$ complex, and can bind laminin 1 even without stimulation (Niessen et al., 1994). We found that these cells bound to both laminin 1 and laminin 8 without stimulation, although activation of the $\beta 1$ integrins with PMA resulted in increased adhesion. The adhesion of non-stimulated cells could be completely inhibited with anti-integrin $\alpha 6$, but only partially with anti- $\beta 1$, again indicating that $\alpha 6\beta 4$ is able to mediate adhesion to r-laminin 8 (Figure 7). In contrast, K562 cells expressing $\alpha 3\beta 1$ adhered poorly to both laminin isoforms (not shown). This agrees with an earlier study where $\alpha 3$ -transfected K562 cells were found to bind efficiently to laminin 8, but poorly to laminin 1 (Delwel et al., 1994).

Cell adhesion to both laminin 1 and r-laminin 8 was found to be dependent on divalent cations, since it could be abolished by 5 mM EDTA in all cell lines tested (Figures 4-8). Heparin, when used at 2 mg/ml, had no effect on the adhesion of HT-1080, BCE, and IBE to r-laminin 8 (Figures 4-6). On laminin 1, however, there was a slight decrease in adhesion of BCE cells (Figure 5), while the other cell lines were unaffected (Figures 4,6). The RGDS-peptide that is reported to block the function of various integrins (Pierschbacher and Ruoslahti, 1984, *Nature* 309(5963), 30-3) had no effect at 1 mM concentration on adhesion of HT-1080, BCE, or IBE cells to the laminins (data not shown).

It was further observed that the cell-binding activity of r-laminin 8 was sensitive to air-drying. When the coated protein was allowed to air dry for 15 minutes at room temperature before adding the cells, the cell-binding activity of r-laminin 8 was completely lost (Figure 4). Even shorter than a 15 minute exposure could abolish the cell-binding activity (not shown). A drop of buffer was allowed to sit on the plastic, while the rest of the well was briefly exposed to air drying. On the dried area, the BCE cells were rounded, and only a few of them showed any signs of spreading. On the area kept wet, practically all cells were well spread and tightly adhered to the surface. Accordingly, all cells on the dried area were lost during washing. Laminin 1 was not as sensitive to this effect, but drying still reduced the cell binding activity by half (Figure 4).

DISCUSSION

The present work provides significant advances concerning the recently described laminin 8 isoform and its $\alpha 4$ chain. Large quantities of r-laminin 8 could be produced as native trimeric protein in cultured human cells, and the r-laminin 8 was shown to be biologically active and to have cell adhesive properties. Furthermore, r-laminin 8 was shown to have a preference for binding to the $\alpha 6$ integrins.

The r-laminin 8 produced in this study is a species hybrid of two human ($\alpha 4$ and $\gamma 1$) and one mouse ($\beta 1$) chains, and it contained a FLAG epitope tag attached to the C-terminus of the $\alpha 4$ chain. Despite these modifications, r-laminin 8 assembled into trimers in a manner expected from a native laminin protein, as demonstrated by rotary shadowing EM. The amount of r-laminin 8 produced by the HEK-293 cells in

monolayer cultures was quite high considering the size and complexity of the protein. An amount of 3-5 mg/L of culture medium is similar to what is frequently obtained in eukaryotic systems, such as the baculovirus insect cell system.

Similarly to other laminin isoforms characterized to date, all the chains of the r-laminin 8 trimer were disulfide linked to each other. Only a minor fraction consisted of disulfide-linked $\beta 1/\gamma 1$ containing dimers and non-crosslinked $\alpha 4$. These chains were also associated into trimers, since the dimers followed the FLAG-tagged $\alpha 4$ chain in immunoprecipitations using the anti-FLAG mAb. The presence of the $\alpha 4$ chain in r-laminin 8 trimers was also demonstrated by showing that all of the $\alpha 4$ could be immunoprecipitated after several rounds of immunoprecipitation with the anti-laminin 1 Ab that recognizes the $\alpha 1$, $\beta 1$, and $\gamma 1$ chains (data not shown).

The reason for the two minor r-laminin 8 bands of different size reacting with EHS and $\gamma 1$ antibodies is unclear. The larger one agrees with the size for a dimer, but the smaller one could not be accounted for. The size difference could be as large as 100 kD. It is possible that these dimers and non-covalent trimers are the products of incomplete or incorrect post-translational processing due to overexpression.

The purified r-laminin 8 was shown to have biological activity, as all cell lines tested in this study adhered to and spread equally well on r-laminin 8 as on laminin 1. This activity could be abolished by drying the protein, suggesting that native conformation was important for full cell binding activity. The cell binding in all cases be abolished by EDTA, indicating dependence on divalent cations.

A large variety of integrins have been implicated as receptors for different laminin isoforms. In this study, we demonstrated that integrins $\alpha 6\beta 1$ and $\alpha 6\beta 4$ were major mediators of cell adhesion to r-laminin 8. The adhesion of HT-1080 and BCE cells was completely blocked by anti-integrin $\alpha 6$ mAb, despite the fact that both cell lines express a wide spectrum of $\beta 1$ and αv integrins, including several of those shown to bind to other laminin isoforms. (Conforti et. al., 1994, *Cell Adhes. Commun.* 1(4), 279-93) HT-1080 cell adhesion to r-laminin 8 is mediated solely by integrin $\alpha 6\beta 1$, since the adhesion could be blocked not only by anti- $\alpha 6$ mAb, but also by the $\beta 1$ antibody. In contrast, the $\beta 1$ mAb only partially blocked adhesion to BCE cells, suggesting that $\alpha 6\beta 4$ contributed to the binding of BCE cells to r-laminin 8. The role

of $\alpha 6 \beta 4$ as a r-laminin 8 receptor was confirmed by assaying the binding of $\alpha 6$ and $\beta 4$ transfected K562 cells that express both $\alpha 6 \beta 1$ and $\alpha 6 \beta 4$ on the cell surface. Indeed, adhesion was completely blocked with $\alpha 6$ mAb, but only partially with $\beta 1$ mAb, indicating that the $\alpha 6 \beta 4$ complex also binds to r-laminin 8. K562 cells expressing
5 $\alpha 6 \beta 1$ bound r-laminin 8 while $\alpha 3 \beta 1$ expressing cells did not, thus confirming that integrin $\alpha 6 \beta 1$ binds r-laminin 8. Our results somewhat contradict the reported lack of integrin $\alpha 6$ subunit in IBE cells (Kanda et al, 1999), since the adhesion to r-laminin 8 was severely perturbed by the anti-integrin $\alpha 6$ mAb. The result suggests that these cells use yet another receptor(s) in addition to $\alpha 6$ integrins for binding to r-laminin 8.
10 However, in certain cases GoH3 is not able to completely block integrin $\alpha 6$ in $\alpha 6 \beta 4$ complexes (Sonnenberg et al., 1993, *J. Cell Sci.* 106(Pt 4), 1083-102). Thus, the remaining adhesion could be due to incompletely blocked $\alpha 6 \beta 4$ complexes.

Interestingly, adhesion of the cell lines tested to r-laminin 8 was found to be more dependent on integrin $\alpha 6$ than adhesion of the cell lines to laminin 1. Another
15 indication of the different adhesive properties of r-laminin 8 and laminin 1 was the finding that $\alpha 6 \beta 1$ -expressing K562 cells did bind to r-laminin 8 without stimulation, but, as also previously reported (Delwel et al., 1993), needed to be stimulated by PMA to efficiently bind to laminin 1 coated surfaces. Thus, r-laminin 8 appears to have a higher avidity or affinity than laminin 1 to $\alpha 6 \beta 1$. The $\alpha 6 \beta 1$ integrin might bind r-
20 laminin 8 even in the conformation that makes it unable to bind to laminin 1, or the cells could be stimulated by the presence of r-laminin 8 via an unknown mechanism. It could be that the avidity/affinity difference is of biological significance, and may well be one reason for the existence of large numbers of laminin isoforms.

In addition to integrins, several other cell surface proteins have been reported to
25 function as laminin receptors. Alpha-dystroglycan is a component of the dystrophin-dystroglycan complex in the skeletal muscle thought to connect the contractile cytoskeleton to the extracellular matrix. Dystroglycan has also been shown to bind laminin 2 and dystrophin, forming a link between the two. (Ervasti and Campbell, 1993, *J. Cell Biol.* 122(4), 809-23) Indirect evidence suggests that laminin 8 might
30 bind to α -dystroglycan; it has been shown that laminin from laminin α d-deficient dystrophic muscle bound dystroglycan, but, in contrast to laminin from normal muscle,

in a manner that was sensitive to inhibition by heparin. (McDearmon et al., 1998, *J. Biol. Chem.* 273(37), 24139-44) Since upregulation of laminin $\alpha 4$ has been observed in laminin $\alpha 2$ deficient muscular dystrophy (Patton et al, 1997; Ringelmann et al., 1999), it can be assumed that the laminin $\alpha 4$ chain is involved in the observed interaction. Alpha-dystroglycan is not restricted to skeletal muscle. (Durbeej et al. 1998, *J. Histochem. Cytochem.* 46(4), 449-57) It was recently shown to be a receptor for laminin 1 in bovine aorta endothelial cells, binding in a manner sensitive to heparin, dextran sulfate, and fucoidan. (Shimizu et al., 1999, *J. Biol. Chem.* 274(17), 11995-2000) Heparin-sensitive interactions were not detected in this study, but this does not rule out the possibility of such interactions in other cell types or in vivo. We did observe that the r-laminin 8 binds heparin-sepharose at physiological salt concentration (data not shown).

In this study, integrins $\alpha 6\beta 1$ and $\alpha 6\beta 4$ were identified as receptors for r-laminin 8 in cultured cells, and thus it is likely that these integrins mediate binding of laminin 8 in vivo, such as to endothelial and muscle cells. Endothelial cells express a wide variety of integrins depending on developmental stage, activation state, and location. At least integrins $\alpha 2\beta 1$, $\alpha 5\beta 1$, $\alpha 6\beta 1$, $\alpha 6\beta 4$, and $\alpha v\beta 3$ have been found in endothelial cells in vivo (Sonnenberg 1990; Conforti 1992), whereas the main laminin isoforms in endothelial basement membranes (BM) are laminins 8 and 10. Other cells besides endothelial cells are likely to interact with the laminin 8 in endothelial BM; platelets contain and secrete laminin 8 when stimulated and adhere to it using the $\alpha 6\beta 1$ integrin.

Laminins 8 and 9 are also found in developing muscle and in the peripheral nervous system, overlapping in expression with integrin $\alpha 6$. In laminin $\alpha 2$ -deficient muscle, both the laminin $\alpha 4$ and integrin $\alpha 6$ are upregulated. (Vachon et al, 1997, *J. Clin. Invest.* 100(7), 1870-81) Interestingly, integrin $\alpha 6$ and integrin $\beta 4$ knock-outs result in epidermolysis bullosa (Georges-Labouesse et al, 1996, *Nat. Genet.* 13(3), 370-3; van der Neut et al., 1996, *Nat. Genet.* 13(3), 366-9), but no muscular or vascular phenotype was reported.

The present invention is not limited by the aforementioned particular preferred embodiments. It will occur to those ordinarily skilled in the art that various modifications may be made to the disclosed preferred embodiments without diverting

from the concept of the invention. All such modifications are intended to be within the scope of the present invention.

We claim

1. Substantially purified laminin 8.
2. The substantially purified laminin 8 of claim 1, comprising recombinant laminin
5 8.
3. The substantially purified recombinant laminin 8 of claim 2 comprising:
a first chain comprising a polypeptide that is substantially similar to an $\alpha 4$
laminin chain;
a second chain comprising a polypeptide that is substantially similar to a $\beta 1$
10 laminin chain; and
a third chain comprising a polypeptide that is substantially similar to a $\gamma 1$
laminin chain;
wherein the first, second, and third chains are assembled into recombinant
heterotrimeric laminin 8.
15
4. The substantially purified recombinant laminin 8 of claim 2 comprising:
a first chain encoded by a polynucleotide that hybridizes under high stringency
conditions to a coding region of one or more of SEQ ID NO:1, 3, 5, 7, 9, or fragments
thereof;
20 a second chain encoded by a polynucleotide that hybridizes under high
stringency conditions to a coding region of one or more of SEQ ID NO:11, 13, 15, 17,
or fragments thereof; and
a third chain encoded by a polynucleotide that hybridizes under high stringency
conditions to a coding region of one or more of SEQ ID NO: 19, 21, 23, 25, or
25 fragments thereof;
wherein the first, second, and third chains are assembled into recombinant
heterotrimeric laminin 8.
5. The substantially purified recombinant laminin 8 of claim 2 comprising:
30 a first chain comprising a polypeptide at least 70% identical to one or more of
SEQ ID NO:2, 4, 6, 8, 10 or fragments thereof;
a second chain comprising a polypeptide at least 70% identical to one or more

of SEQ ID NO:12, 14, 16, 18 or fragments thereof; and

a third chain comprising a polypeptide at least 70% identical to one or more of SEQ ID NO:20, 22, 24, 26, or fragments thereof;

wherein the first, second, and third chains are assembled into recombinant
5 heterotrimeric laminin 8.

6. The substantially purified recombinant laminin 8 of claim 2 comprising a first, second, and third polypeptide chain, wherein the first, second, and third polypeptide chains each comprise a general structure selected from the group consisting of: (1) R1-
10 R2-R3; (2) R1-R2-R3(e); (3) R3; (4) R3(e); (5) R1-R3; (6) R1-R3(e); (7) R2-R3; and (8) R2-R3(e)

wherein R1 is a amino terminal methionine; R2 is a signal sequence that is capable of directing secretion of the polypeptide, wherein the signal sequence may be the natural signal sequence for the particular laminin chain, that of another secreted
15 protein, or it may be an artificial sequence; R3 is a secreted $\alpha 4$ laminin chain for the first polypeptide chain, a secreted $\beta 1$ laminin chain for the second polypeptide chain, and $\gamma 1$ laminin chain for the third polypeptide chain; and R3(e) is identical to R3, but further comprises an epitope tag.

20 7. Recombinant laminin 8-expressing host cells.

8. The recombinant laminin 8-expressing host cells of claim 7, wherein the cells express recombinant laminin 8 comprising:

a first chain comprising a recombinant polypeptide that is substantially similar
25 to an $\alpha 4$ laminin polypeptide;

a second chain comprising a recombinant polypeptide that is substantially similar to a $\beta 1$ laminin polypeptide sequence; and

a third chain comprising a recombinant polypeptide that is substantially similar to a $\gamma 1$ laminin polypeptide sequence;

30 wherein the cell expresses the first, second, and third chains, and wherein the first, second, and third chains assemble into recombinant laminin 8 that is secreted into the media by the cultured cell.

9. The recombinant laminin 8-expressing host cells of claim 7, wherein the cells express recombinant laminin 8 comprising:

a first chain encoded by a polypeptide that hybridizes under high stringency conditions to a coding region of one or more of SEQ ID NO:1, 3, 5, 7, or 9, or fragments thereof;

a second chain encoded by a polypeptide that hybridizes under high stringency conditions to a coding region of one or more of SEQ ID NO:11, 13, or fragments thereof; and

a third chain encoded by a polypeptide that hybridizes under high stringency conditions to a coding region of one or more of SEQ ID NO: 15, 17, or fragments thereof;

wherein the cell expresses the first, second, and third chains, and wherein the first, second, and third chains assemble into recombinant laminin 8 that is secreted into the media by the cultured cell.

10. The recombinant laminin 8-expressing host cells of claim 7, wherein the cells express recombinant laminin 8 comprising:

a first chain comprising a polypeptide at least 70% identical to one or more of SEQ ID NO:2, 4, 6, 8, 10 or fragments thereof;

a second chain comprising a polypeptide at least 70% identical to one or more of SEQ ID NO:12, 14, 16, 18 or fragments thereof; and

a third chain comprising a recombinant polypeptide at least 70% identical to one or more of SEQ ID NO:20, 22, 24, 26, or fragments thereof;

wherein the cell expresses the first, second, and third chains, and wherein the first, second, and third chains assemble into recombinant laminin 8 that is secreted into the media by the cultured cell.

11. The recombinant laminin 8-expressing host cells of claim 7, wherein the cells express recombinant laminin 8 comprising a first, second, and third polypeptide chain, wherein the first, second, and third polypeptide chains each comprise a general

structure selected from the group consisting of: (1) R1-R2-R3; (2) R1-R2-R3(e); (3) R3; (4) R3(e); (5) R1-R3; (6) R1-R3(e); (7) R2-R3; and (8) R2-R3(e)

wherein R1 is a amino terminal methionine; R2 is a signal sequence that is capable of directing secretion of the polypeptide, wherein the signal sequence may be the natural signal sequence for the particular laminin chain, that of another secreted protein, or it may be an artificial sequence; R3 is a secreted $\alpha 4$ laminin chain for the first polypeptide chain, a secreted $\beta 1$ laminin chain for the second polypeptide chain, and $\gamma 1$ laminin chain for the third polypeptide chain; and R3(e) is identical to R3, but further comprises an epitope tag .

10

12. The host cells of any of claims 7-11, wherein the host cell is a mammalian cell.

13. The host cells of claim 12, wherein at least one of the first, second, or third chains is expressed as a fusion protein with an epitope tag.

15

14. A method of purifying recombinant laminin 8, comprising:

a. providing the host cells of claim 12;

b. growing the cells in cell culture medium under conditions to stimulate expression of the recombinant laminin 8 chains;

20

c. passing the cell culture medium through an affinity chromatography column, wherein the column contains a compound that binds to the recombinant laminin 8;

25

d. washing the affinity column to remove unbound materials; and

e. eluting the bound recombinant laminin 8 from the column.

15. Substantially purified recombinant laminin 8 isolated according to the method of claim 14.

30

16. A pharmaceutical composition comprising:
- a. laminin 8; and
 - b. a pharmaceutically acceptable carrier.
- 5 17. The pharmaceutical composition of claim 16, wherein the laminin 8 comprises recombinant laminin 8.
18. A method to accelerate healing of a vascular tissue injury in a subject, comprising contacting the site of the vascular tissue injury of the subject with an
10 amount effective of laminin 8 to promote re-endothelialization at the vascular tissue injury site.
19. The method of claim 18, wherein the vascular injury is selected from the group consisting of angioplasty restenosis, vascular surgical procedures, aneurysm, and
15 atherosclerosis.
20. A method to accelerate healing of a bone or connective tissue injury in a subject comprising contacting the site of the bone or connective tissue injury in the subject with an amount effective of laminin 8 to accelerate healing of the bone or connective tissue
20 injury.
21. The method of claim 20 wherein healing is accomplished by incorporation of recombinant laminin 8 into wound repair dressings, matrices, or tissue grafts.
- 25 22. The method of claim 20 wherein the bone or connective tissue injury is selected from the group consisting of fractures, tears, deformities, or defects of bone, tendon, cartilage, and ligament.
23. A method to improve the biocompatibility of a medical device or graft,
30 comprising contacting the medical device or graft with an amount effective of laminin 8 to improve the biocompatibility of the medical device or graft.

24. An improved medical device or graft, wherein the improvement consists of providing a medical device or graft with an amount effective of laminin 8 to improve the biocompatibility of the medical device or graft.

5 25. A method to regulate angiogenesis in a subject, comprising contacting a site in need of angiogenesis in the subject with an amount effective of laminin 8 to regulate angiogenesis.

10 26. A method to promote neural regeneration in a subject, comprising contacting a site in need of neural regeneration in the subject with an amount effective of laminin 8 to promote neural regeneration.

27. A method to promote cell adhesion to a surface, comprising contacting cells with an amount effective of the laminin 8 to promote cell adhesion to the surface.

15

28. An improved cell growth substrate, wherein the improvement consists of providing a cell growth substrate that has been coated with an amount effective of laminin 8 to promote cell attachment to the cell growth substrate.

20 29. The method of any of claims 18-28, wherein the laminin 8 comprises recombinant laminin 8.

30. A method to accelerate healing of a vascular tissue injury in a subject, comprising contacting the site of the vascular tissue injury of the subject with an amount effective of the pharmaceutical composition of claim 16 or 17 to promote re-endothelialization at the vascular tissue injury site.

31. The method of claim 30, wherein the vascular injury is selected from the group consisting of angioplasty restenosis, vascular surgical procedures, aneurysm, and atherosclerosis

30

32. A method to accelerate healing of a bone or connective tissue injury in a subject comprising contacting the site of the bone or connective tissue injury in the subject with an amount effective of the pharmaceutical composition of claim 16 or 17 to accelerate healing of the bone or connective tissue injury.

5

33. The method of claim 32 wherein healing is accomplished by incorporation of recombinant laminin 8 into wound repair dressings, matrices, or tissue grafts.

34. The method of claim 32 wherein the bone or connective tissue injury is selected
10 from the group consisting of fractures, tears, deformities, or defects of bone, tendon, cartilage, and ligament.

35. A method to improve the biocompatibility of a medical device or graft, comprising contacting the medical device or graft with an amount effective of the
15 pharmaceutical composition of claim 16 or 17 to improve the biocompatibility of the medical device or graft.

36. An improved medical device or graft, wherein the improvement consists of providing a medical device or graft with an amount effective of the pharmaceutical
20 composition of claim 16 or 17 to improve the biocompatibility of the medical device or graft.

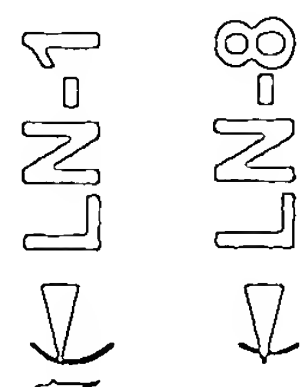
37. A method to promote angiogenesis in a subject, comprising contacting a site in need of angiogenesis in the subject with an amount effective of the pharmaceutical
25 composition of claim 16 or 17 to promote angiogenesis.

38. A method to promote neural regeneration in a subject, comprising contacting a site in need of neural regeneration in the subject with an amount effective of the pharmaceutical composition of claim 16 or 17 to promote neural regeneration.

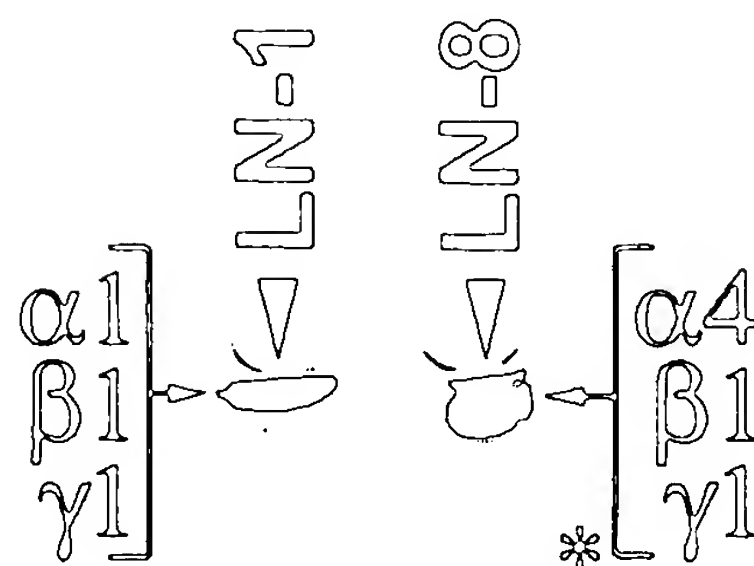
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39. A method to promote cell adhesion to a surface, comprising contacting cells with an amount effective of the pharmaceutical composition of claim 16 or 17 to promote cell adhesion to the surface.
- 5 40. An improved cell growth substrate, wherein the improvement consists of providing a cell growth substrate that has been coated with an amount effective of the pharmaceutical composition of claim 16 or 17 to promote cell attachment to the cell growth substrate.
- 10 41. A kit for carrying out the method of any of claims 18-28, comprising:
(a) an amount effective of laminin 8 for carrying out the method; and
(b) instructions for using the laminin 8 for carrying out the method.
- 15 42. A method to inhibit cell adhesion to laminin 8, comprising contacting the cell with an amount effective of an antagonist of at least one of integrin $\alpha 6 \beta 1$ and $\alpha 6 \beta 4$ to inhibit cell adhesion to laminin 8.

reduced

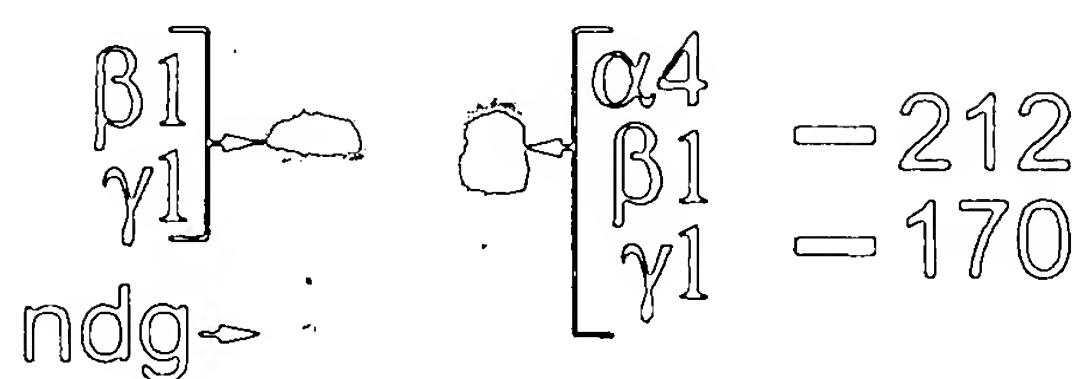


unreduced



$\alpha 1 \rightarrow$

$\rightarrow *$



$\text{ndg} \rightarrow$

$\rightarrow \alpha 4$

$= 116$

$= 76$

$= 53$



FIG. 1

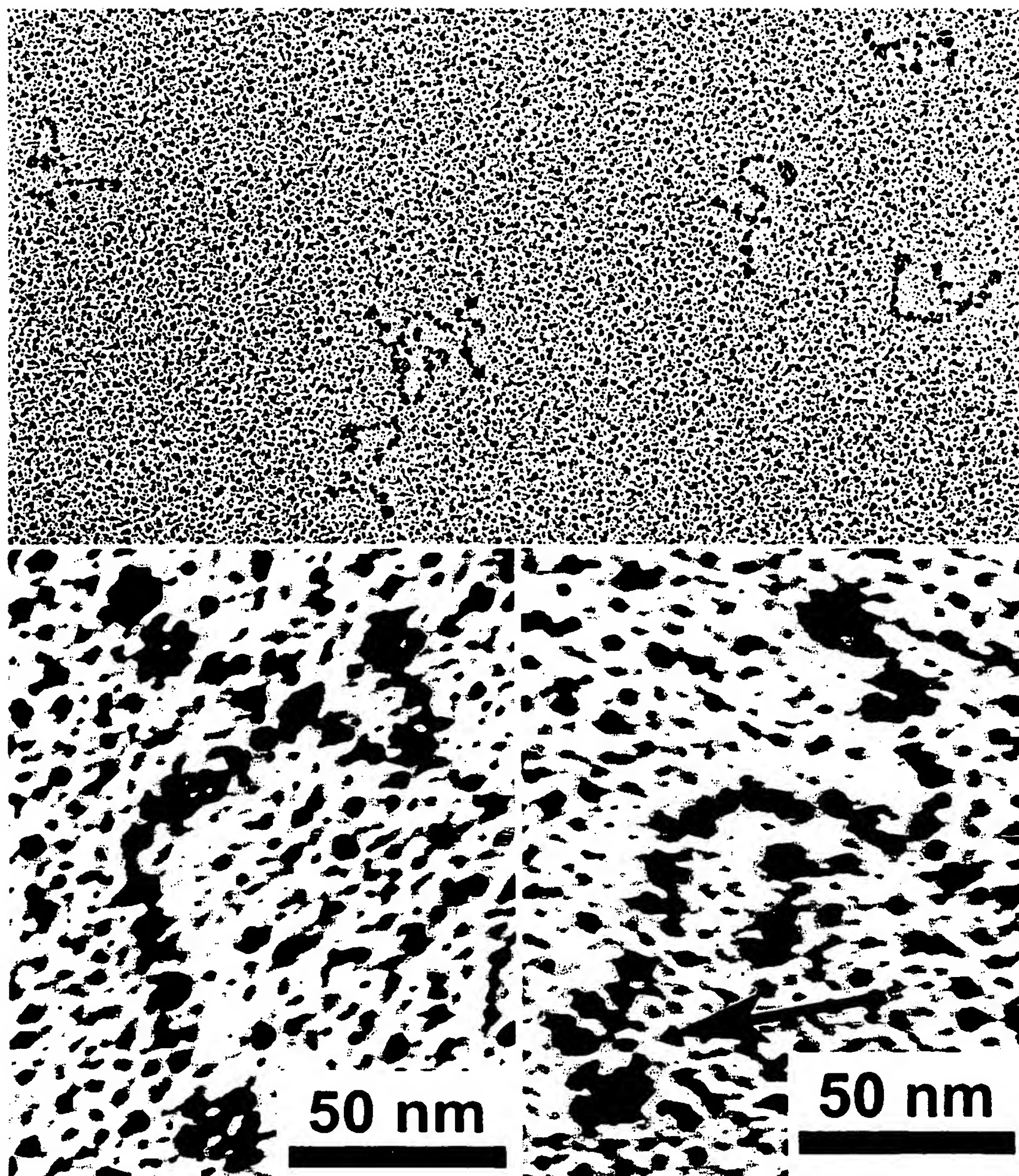
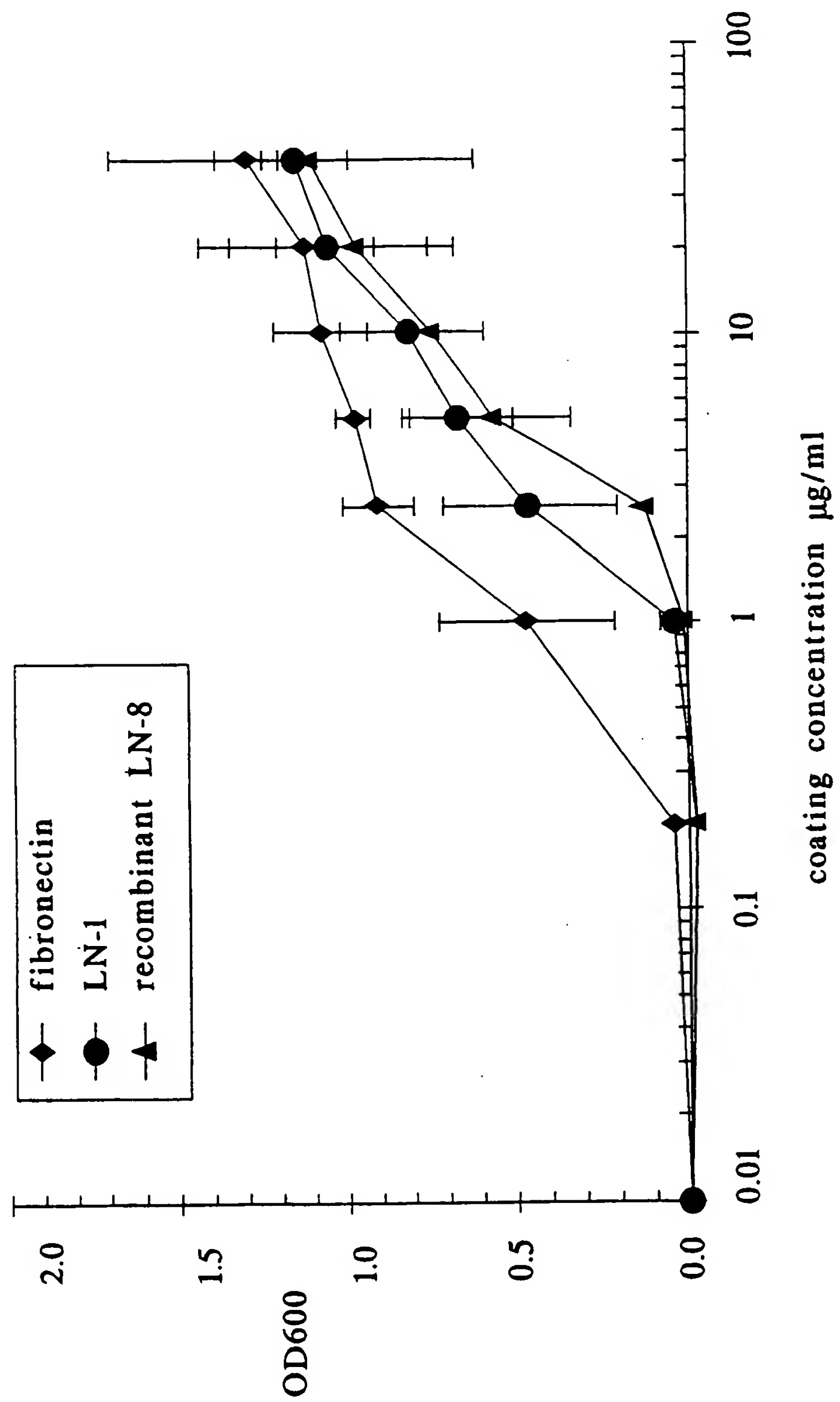
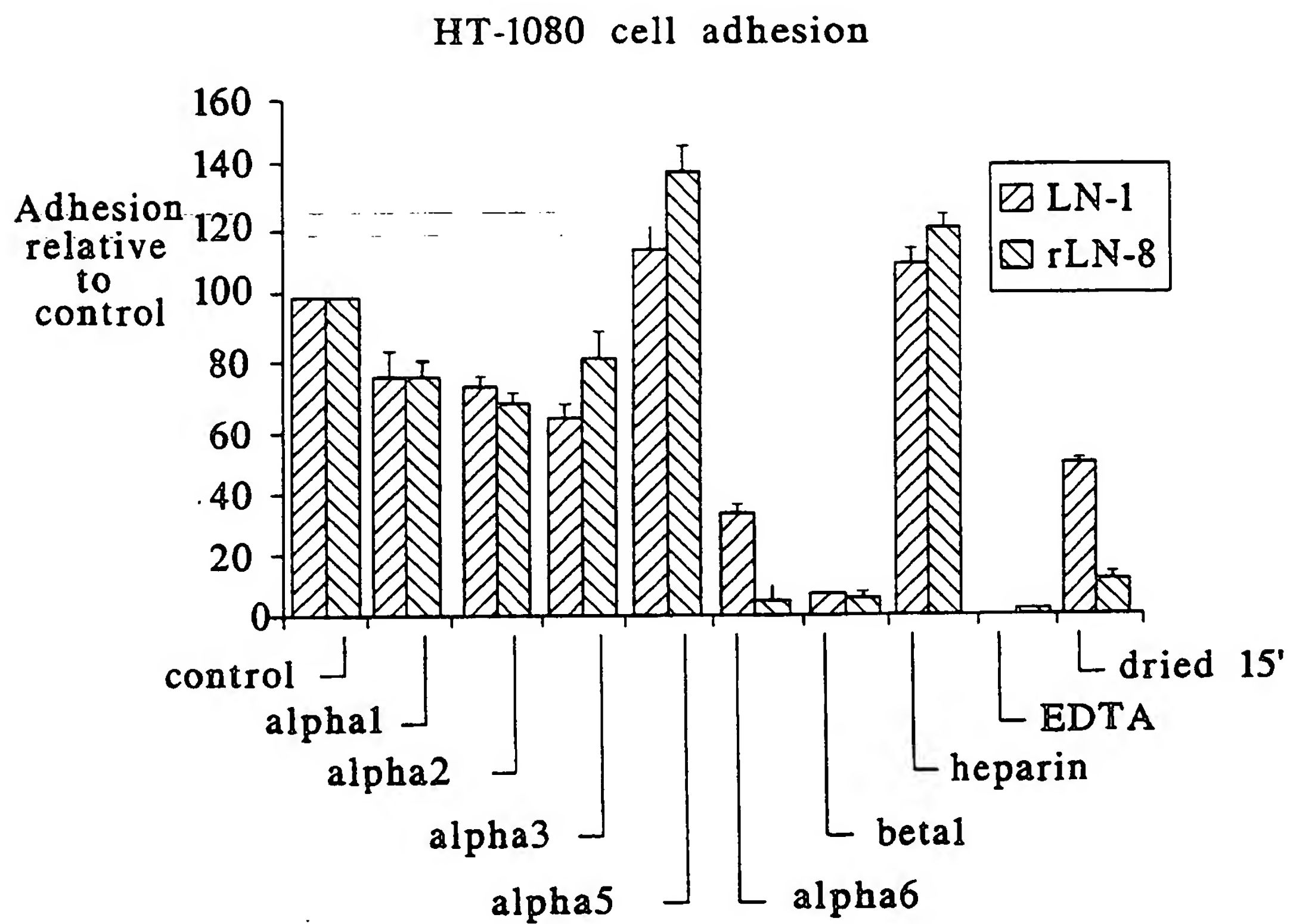


FIG. 2

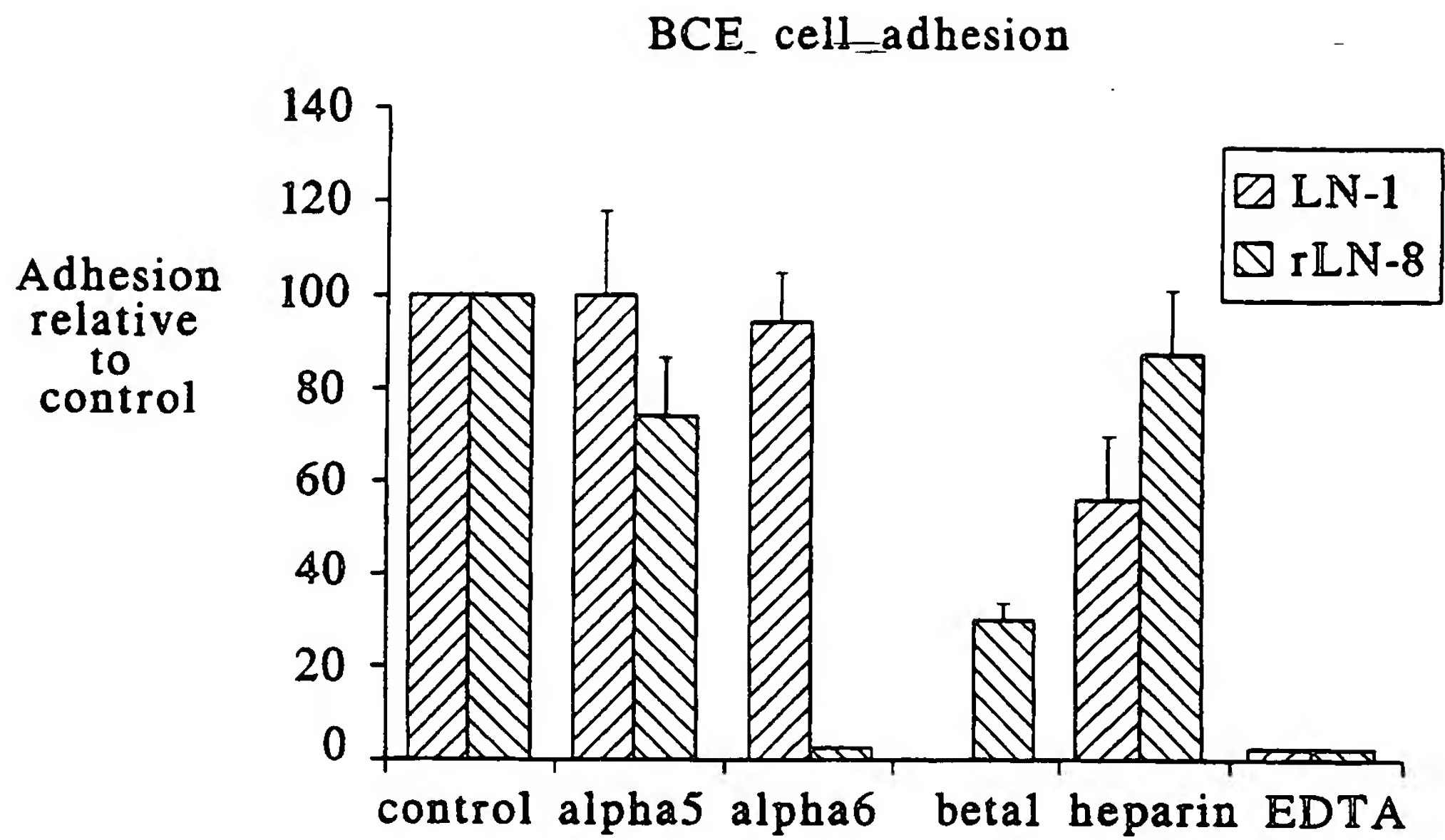
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FIG. 3

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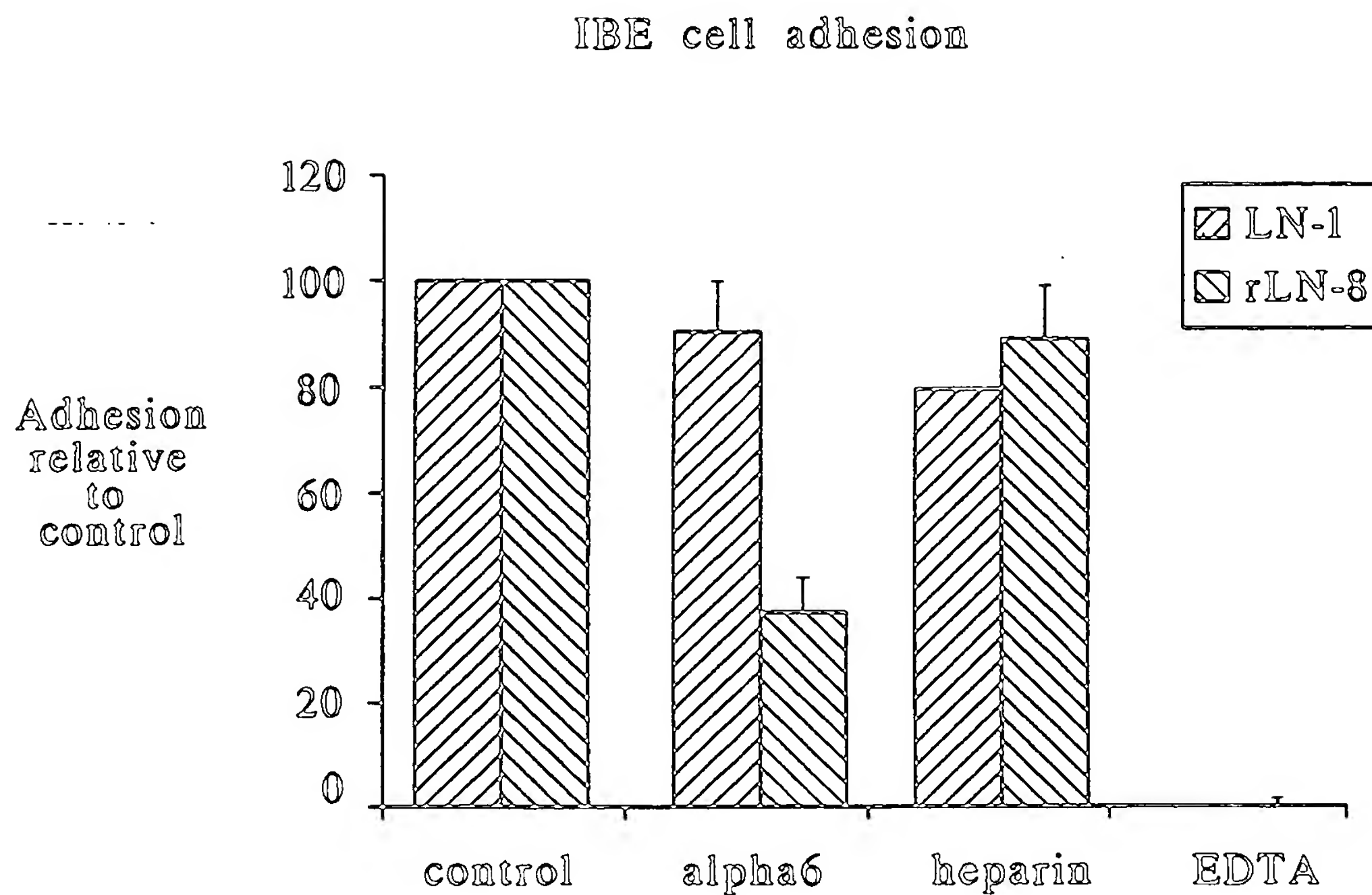
FIG. 4

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FIG. 5

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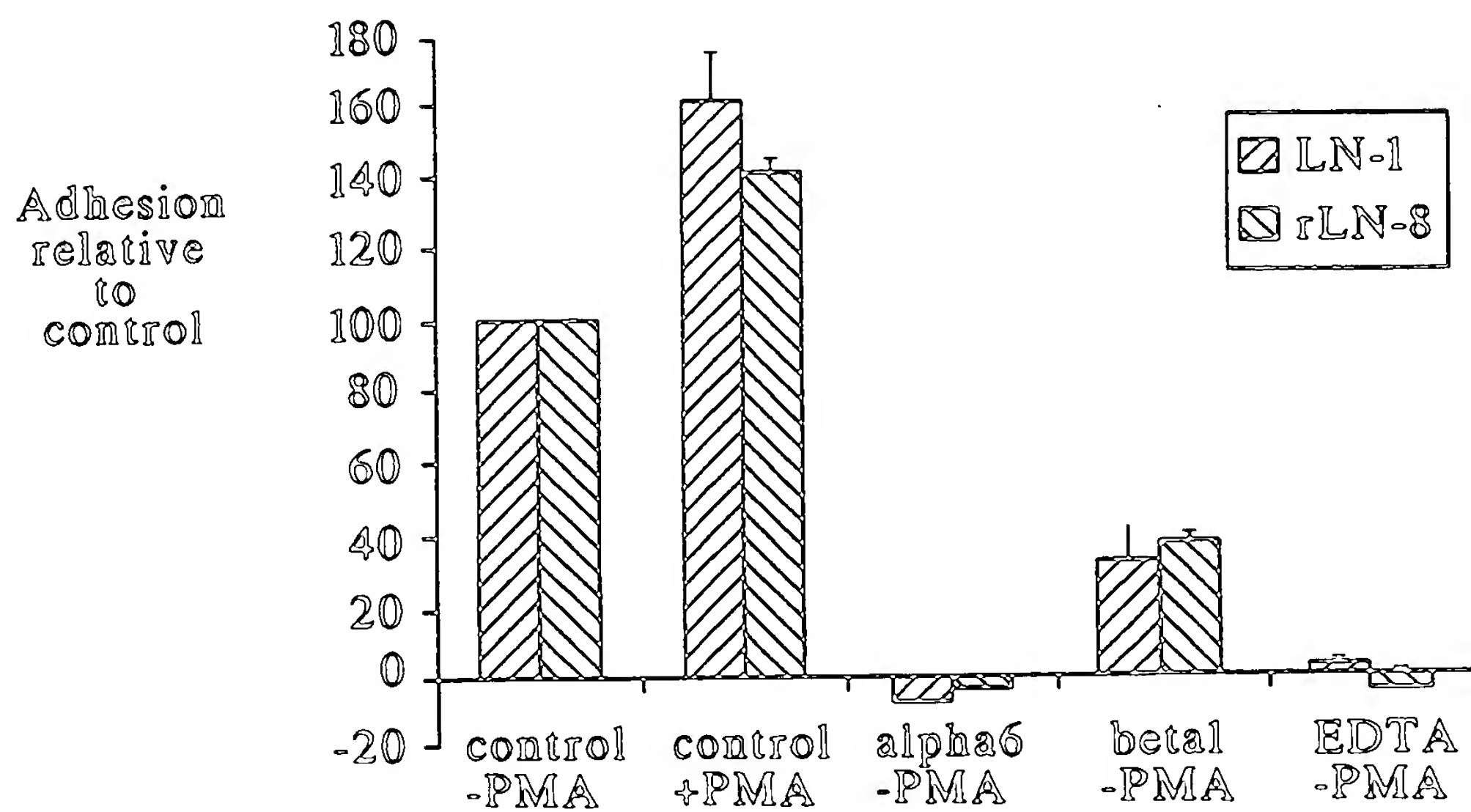
FIG. 6



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FIG. 7

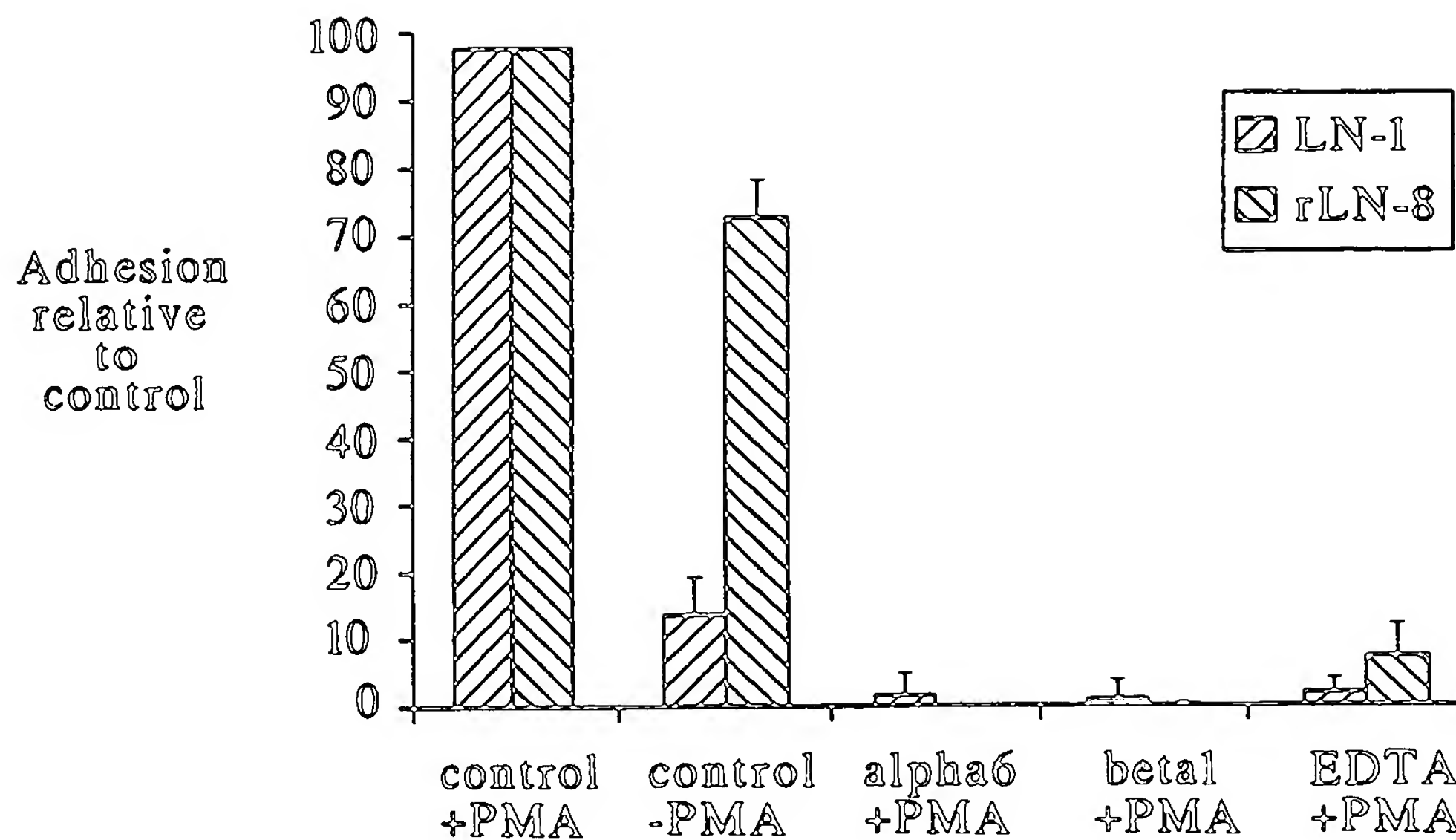
K562 Alpha6Beta4/Alpha6Beta1



8/8

FIG. 8

K562 Alpha6Beta1



SEQUENCE LISTING

<110> Korttesmaa, Jarrko
Tryggvason, Karl

<120> Laminin 8 and Methods For Its Use

<130> 99,274-D1

<140> To Be Assigned

<141> Filed Herewith

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<170> PatentIn Ver. 2.0

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ata gat gga gcc aaa agt gaa cta caa gta aaa cta tct aac cta agt Ile Asp Gly Ala Lys Ser Glu Leu Gln Val Lys Leu Ser Asn Leu Ser 560 565 570			1909
aac ctc agc cat gat tta gtc caa gaa gct att gac cat gca cag gac Asn Leu Ser His Asp Leu Val Gln Glu Ala Ile Asp His Ala Gln Asp 575 580 585			1957

ctt	caa	caa	gaa	gct	aat	gaa	ttg	agc	agg	aag	ttg	cac	agt	tca	gat	2005
Leu	Gln	Gln	Glu	Ala	Asn	Glu	Leu	Ser	Arg	Lys	Leu	His	Ser	Ser	Asp	
590					595					600					605	
atg	aac	ggg	ctg	gta	cag	aag	gct	ttg	gat	gca	tca	aat	gtc	tat	gaa	2053
Met	Asn	Gly	Leu	Val	Gln	Lys	Ala	Leu	Asp	Ala	Ser	Asn	Val	Tyr	Glu	
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aat	att	gtt	aat	tat	gtt	agt	gaa	gcc	aat	gaa	aca	gca	gaa	ttt	gct	2101
Asn	Ile	Val	Asn	Tyr	Val	Ser	Glu	Ala	Asn	Glu	Thr	Ala	Glu	Phe	Ala	
			625					630						635		
ttg	aac	acc	act	gac	cga	att	tat	gat	gcg	gtg	agt	ggg	att	gat	act	2149
Leu	Asn	Thr	Thr	Asp	Arg	Ile	Tyr	Asp	Ala	Val	Ser	Gly	Ile	Asp	Thr	
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caa	atc	att	tac	cat	aaa	gat	gaa	agt	gag	aac	ctc	ctc	aat	caa	gcc	2197
Gln	Ile	Ile	Tyr	His	Lys	Asp	Glu	Ser	Glu	Asn	Leu	Leu	Asn	Gln	Ala	
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Arg	Glu	Leu	Gln	Ala	Lys	Ala	Glu	Ser	Ser	Ser	Asp	Glu	Ala	Val	Ala	
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Asp	Thr	Ser	Arg	Arg	Val	Gly	Gly	Ala	Leu	Ala	Arg	Lys	Ser	Ala	Leu	
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Lys	Thr	Arg	Leu	Ser	Asp	Ala	Val	Lys	Gln	Leu	Gln	Ala	Ala	Glu	Arg	
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ggg	gat	gcc	cag	cag	cgc	ctg	ggg	cag	tct	aga	ctg	atc	acc	gag	gaa	2389
Gly	Asp	Ala	Gln	Gln	Arg	Leu	Gly	Gln	Ser	Arg	Leu	Ile	Thr	Glu	Glu	
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gcc	aac	agg	acg	acg	atg	gag	gtg	cag	cag	gcc	act	gcc	ccc	atg	gcc	2437
Ala	Asn	Arg	Thr	Thr	Met	Glu	Val	Gln	Gln	Ala	Thr	Ala	Pro	Met	Ala	
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aac	aat	cta	acc	aac	tgg	tca	cag	aat	ctt	caa	cat	ttt	gac	tct	tct	2485
Asn	Asn	Leu	Thr	Asn	Trp	Ser	Gln	Asn	Leu	Gln	His	Phe	Asp	Ser	Ser	
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gct	tac	aac	act	gca	gtg	aac	tct	gct	agg	gat	gca	gta	aga	aat	ctg	2533
Ala	Tyr	Asn	Thr	Ala	Val	Asn	Ser	Ala	Arg	Asp	Ala	Val	Arg	Asn	Leu	
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acc	gag	gtt	gtc	cct	cag	ctc	ctg	gat	cag	ctt	cgt	acg	gtt	gag	cag	2581
Thr	Glu	Val	Val	Pro	Gln	Leu	Leu	Asp	Gln	Leu	Arg	Thr	Val	Glu	Gln	
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aag	cga	cct	gca	agc	aac	gtt	tct	gcc	agc	atc	cag	agg	atc	cga	gag	2629
Lys	Arg	Pro	Ala	Ser	Asn	Val	Ser	Ala	Ser	Il	Gln	Arg	Ile	Arg	Glu	
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ctc	att	gct	cag	acc	aga	agt	gtt	gcc	agc	aag	atc	caa	gtc	tcc	atg	2677
Leu	Ile	Ala	Gln	Thr	Arg	Ser	Val	Ala	Ser	Lys	Ile	Gln	Val	Ser	Met	
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Asp Asp Leu Lys Ala Phe Thr Ser Leu Ser Leu Tyr Met Lys Pro Pro	
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Val Lys Arg Pro Glu Leu Thr Glu Thr Ala Asp Gln Phe Ile Leu Tyr	
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Leu Gly Ser Lys Asn Ala Lys Lys Glu Tyr Met Gly Leu Ala Ile Lys	
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Asn Asp Asn Leu Val Tyr Val Tyr Asn Leu Gly Thr Lys Asp Val Glu	
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Ile Val Lys Ile Glu Arg Val Gly Lys His Gly Lys Val Phe Leu Thr	
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Val Pro Ser Leu Ser Ser Thr Ala Glu Glu Lys Phe Ile Lys Lys Gly	
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Ser Glu Lys Lys Phe Tyr Phe Gly Gly Ser Pro Ile Ser Ala Gln Tyr				
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Arg Asp Val Glu Val Glu Asp Phe Gln Arg Tyr Thr Glu Lys Val His				
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Thr Ser Leu Tyr Glu Cys Pro Ile Glu Ser Ser Pro Leu Phe Leu Leu				
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His Lys Lys Gly Lys Asn Leu Ser Lys Pro Lys Ala Ser Gln Asn Lys				
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Phe Tyr Val Ser Asp Gln Glu Glu Asn Asp Phe Met Thr Leu Phe Leu				
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Ala His Gly Arg Leu Val Tyr Met Phe Asn Val Gly His Lys Lys Leu				
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Ile Phe Ile Arg Glu Arg Ser Ser Gly Arg Leu Val Ile Asp Gly Leu				
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Arg Val Leu Glu Glu Ser Leu Pro Pro Thr Glu Ala Thr Trp Lys Ile	
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His Arg Glu Pro Val Phe Val Gly Gly Val Pro Glu Ser Leu Leu Thr	
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 Glu Thr Ser Glu Pro Arg Val Ala Leu Gly Arg Leu Pro Pro Ala Ala
 50 55 60
 Glu Lys Cys Asn Ala Gly Phe Phe His Thr Leu Ser Gly Glu Cys Val
 65 70 75 80
 Pro Cys Asp Cys Asn Gly Asn Ser Asn Glu Cys Leu Asp Gly Ser Gly
 85 90 95
 Tyr Cys Val His Cys Gln Arg Asn Thr Thr Gly Glu His Cys Glu Lys
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 Cys Leu Asp Gly Tyr Ile Gly Asp Ser Ile Arg Gly Ala Pro Gln Phe
 115 120 125
 Cys Gln Pro Cys Pro Cys Pro Leu Pro His Leu Ala Asn Phe Pro Glu
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 Ser Cys Tyr Arg Lys Asn Gly Ala Val Arg Cys Ile Cys Asn Glu Asn
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Tyr Ala Gly Pro Asn Cys Glu Arg Cys Ala Pro Gly Tyr Tyr Gly Asn
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 Pro Phe Leu Ile Gly Ser Thr Cys Lys Lys Cys Asp Cys Ser Gly Asn
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 Ser Asp Pro Asn Leu Ile Phe Glu Asp Cys Asp Glu Val Thr Gly Gln
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 Cys Arg Asn Cys Leu Arg Asn Thr Thr Gly Phe Lys Cys Glu Arg Cys
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 Ala Pro Gly Tyr Tyr Gly Asp Ala Arg Ile Ala Lys Asn Cys Ala Val
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 Cys Asn Cys Gly Gly Gly Pro Cys Asp Ser Val Thr Gly Glu Cys Leu
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 Ser Gly Val Leu Ser Val Ser Ser Gly Ala Ala Ala His Arg His Val
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Ser His Gly Met Ile Phe Tyr Val Ser Asp Gln Glu Glu Asn Asp Phe	
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Met Thr Leu Phe Leu Ala His Gly Arg Leu Val Tyr Met Phe Asn Val	
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Val Ile Asp Gly Leu Arg Val Leu Glu Glu Ser Leu Pro Pro Thr Glu	
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Ala Thr Trp Lys Ile Lys Gly Pro Ile Tyr Leu Gly Gly Val Ala Pro	
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Gly Lys Ala Val Lys Asn Val Gln Ile Asn Ser Ile Tyr Ser Phe Ser	
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Gly Thr Tyr Phe Ser Thr Glu Gly Gly Tyr Val Val Leu Asp Glu Ser	
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Phe Asn Ile Gly Leu Lys Phe Glu Ile Ala Phe Glu Val Arg Pro Arg
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His Thr Leu Ser Gly Glu Cys Val Pro Cys Asp Cys Asn Gly Asn Ser
 50 55 60

Asn Glu Cys Leu Asp Gly Ser Gly Tyr Cys Val His Cys Gln Arg Asn
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Thr Thr Gly Glu His Cys Glu Lys Cys Leu Asp Gly Tyr Ile Gly Asp
 85 90 95

Ser Ile Arg Gly Ala Pro Gln Phe Cys Gln Pro Cys Pro Cys Pro Leu
 100 105 110

Pro His Leu Ala Asn Phe Pro Glu Ser Cys Tyr Arg Lys Asn Gly Ala
 115 120 125

Val Arg Cys Ile Cys Asn Glu Asn Tyr Ala Gly Pro Asn Cys Glu Arg
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Cys Ala Pro Gly Tyr Tyr Gly Asn Pro Phe Leu Ile Gly Ser Thr Cys
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Lys Lys Cys Asp Cys Ser Gly Asn Ser Asp Pro Asn Leu Ile Phe Glu
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Asp Cys Asp Glu Val Thr Gly Gln Cys Arg Asn Cys Leu Arg Asn Thr
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Thr Gly Phe Lys Cys Glu Arg Cys Ala Pro Gly Tyr Tyr Gly Asp Ala
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Arg Ile Ala Lys Asn Cys Ala Val Cys Asn Cys Gly Gly Gly Pro Cys
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Asp Ser Val Thr Gly Glu Cys Leu Glu Glu Gly Phe Glu Pro Pro Thr
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Gly Cys Asp Lys Cys Val Trp Asp Leu Thr Asp Asp Leu Arg Leu Ala
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Ala Leu Ser Ile Glu Glu Gly Lys Ser Gly Val Leu Ser Val Ser Ser
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Gly Ala Ala Ala His Arg His Val Asn Glu Ile Asn Ala Thr Ile Tyr
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Leu Leu Lys Thr Lys Leu Ser Glu Arg Glu Asn Gln Tyr Ala Leu Arg
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Lys Ile Gln Ile Asn Asn Ala Glu Asn Thr Met Lys Ser Leu Leu Ser
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 Asp Val Glu Glu Leu Val Glu Lys Glu Asn Gln Ala Ser Arg Lys Gly
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 Pro Arg Leu Thr Leu Ser Glu Leu Asp Asp Ile Ile Lys Asn Ala Ser
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 Ser Asn Val Tyr Glu Asn Il Val Asn Tyr Val Ser Glu Ala Asn Glu
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 Thr Ala Glu Phe Ala Leu Asn Thr Thr Asp Arg Ile Tyr Asp Ala Val
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 Pro Ala Tyr Phe Ser Ile Val Lys Ile Glu Arg Val Gly Lys His Gly
 900 905 910
 Lys Val Phe Leu Thr Val Pro Ser Leu Ser Ser Thr Ala Glu Glu Lys
 915 920 925
 Phe Ile Lys Lys Gly Glu Phe Ser Gly Asp Asp Ser Leu Leu Asp Leu
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 Asp Pro Glu Asp Thr Val Phe Tyr Val Gly Gly Val Pro Ser Asn Phe

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	980	985	990
Ile Tyr Asn Met Asp Pro Ser Thr Ser Val Pro Cys Ala Arg Asp Lys			
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Leu Ala Phe Thr Gln Ser Arg Ala Ala Ser Tyr Phe Phe Asp Gly Ser			
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Gly Tyr Ala Val Val Arg Asp Ile Pro Arg Arg Gly Lys Phe Gly Gln			
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Val Thr Arg Phe Asp Ile Glu Val Arg Thr Pro Ala Asp Asn Gly Leu			
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Ile Leu Leu Met Val Asn Gly Ser Met Phe Phe Arg Leu Glu Met Arg			
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Asn Gly Tyr Leu His Val Phe Tyr Asp Phe Gly Phe Ser Ser Gly Arg			
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Val His Leu Glu Asp Thr Leu Lys Lys Ala Gln Ile Asn Asp Ala Lys			
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Tyr His Glu Ile Ser Ile Ile Tyr His Asn Asp Lys Lys Met Ile Leu			
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Val Val Asp Arg Arg His Val Lys Ser Met Asp Asn Glu Lys Met Lys			
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Ile Pro Phe Thr Asp Ile Tyr Ile Gly Gly Ala Pro Pro Glu Ile Leu			
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Lys Gly Ile Lys Val Gln Ser Val Asp Lys Gln Tyr Asn Asp Gly Leu			
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			1280

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Gln Leu Asp Val Asp Ser Glu Val Asn His Val Val Gly Pro Leu Asn
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Glu Leu Leu Ser Gln Ala Glu Ser Trp Gln Arg Leu His Asn Glu Thr	
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735 740 745	
gcc aac aat cta acc aac tgg tca cag aat ctt caa cat ttt gac tct	2308

Ala	Asn	Asn	Leu	Thr	Asn	Trp	Ser	Gln	Asn	Leu	Gln	His	Phe	Asp	Ser		
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Ser	Ala	Tyr	Asn	Thr	Ala	Val	Asn	Ser	Ala	Arg	Asp	Ala	Val	Arg	Asn		
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ctg	acc	gag	gtt	gtc	cct	cag	ctc	ctg	gat	cag	ctt	cgt	acg	gtt	gag	2404	
Leu	Thr	Glu	Val	Val	Pro	Gln	Leu	Leu	Asp	Gln	Leu	Arg	Thr	Val	Glu		
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cag	aag	cga	cct	gca	agc	aac	gtt	tct	gcc	agc	atc	cag	agg	atc	cga	2452	
Gln	Lys	Arg	Pro	Ala	Ser	Asn	Val	Ser	Ala	Ser	Ile	Gln	Arg	Ile	Arg		
			800					805					810				
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Glu	Leu	Ile	Ala	Gln	Thr	Arg	Ser	Val	Ala	Ser	Lys	Ile	Gln	Val	Ser		
		815					820					825					
atg	atg	ttt	gat	ggc	cag	tca	gct	gtg	gaa	gtg	cac	tcg	aga	acc	agt	2548	
Met	Met	Phe	Asp	Gly	Gln	Ser	Ala	Val	Glu	Val	His	Ser	Arg	Thr	Ser		
	830					835					840						
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Met	Asp	Asp	Leu	Lys	Ala	Phe	Thr	Ser	Leu	Ser	Leu	Tyr	Met	Lys	Pro		
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cct	gtg	aag	cgg	ccg	gaa	ctg	acc	gag	act	gca	gat	cag	ttt	atc	ctg	2644	
Pro	Val	Lys	Arg	Pro	Glu	Leu	Thr	Glu	Thr	Ala	Asp	Gln	Phe	Ile	Leu		
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Tyr	Leu	Gly	Ser	Lys	Asn	Ala	Lys	Lys	Glu	Tyr	Met	Gly	Leu	Ala	Ile		
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Lys	Asn	Asp	Asn	Leu	Val	Tyr	Val	Tyr	Asn	Leu	Gly	Thr	Lys	Asp	Val		
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Glu	Ile	Pro	Leu	Asp	Ser	Lys	Pro	Val	Ser	Ser	Trp	Pro	Ala	Tyr	Phe		
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Ser	Ile	Val	Lys	Ile	Glu	Arg	Val	Gly	Lys	His	Gly	Lys	Val	Phe	Leu		
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Gly	Glu	Phe	Ser	Gly	Asp	Asp	Ser	Leu	Leu	Asp	Leu	Asp	Pro	Glu	Asp		
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Asp Pro Ser Thr Ser Val Pro Cys Ala Arg Asp Lys Leu Ala Phe Thr			
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cag agt cgg gct gcc agt tac ttc ttc gat ggc tcc ggt tat gcc gtg			3172
Gln Ser Arg Ala Ala Ser Tyr Phe Phe Asp Gly Ser Gly Tyr Ala Val			
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Val Arg Asp Ile Pro Arg Arg Gly Lys Phe Gly Gln Val Thr Arg Phe			
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gac ata gaa gtt cga aca cca gct gac aac ggc ctt att ctc ctg atg			3268
Asp Ile Glu Val Arg Thr Pro Ala Asp Asn Gly Leu Ile Leu Leu Met			
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Val Asn Gly Ser Met Phe Phe Arg Leu Glu Met Arg Asn Gly Tyr Leu			
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cat gtg ttc tat gat ttt gga ttc agc agt ggc cgt gtg cat ctt gaa			3364
His Val Phe Tyr Asp Phe Gly Phe Ser Ser Gly Arg Val His Leu Glu			
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gat acg tta aag aaa gct caa att aat gat gca aaa tac cat gag atc			3412
Asp Thr Leu Lys Lys Ala Gln Ile Asn Asp Ala Lys Tyr His Glu Ile			
	1120	1125	1130
tca atc att tac cac aat gat aag aaa atg atc ttg gta gtt gac aga			3460
Ser Ile Ile Tyr His Asn Asp Lys Lys Met Ile Leu Val Val Asp Arg			
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agg cat gtc aag agc atg gat aat gaa aag atg aaa ata cct ttt aca			3508
Arg His Val Lys Ser Met Asp Asn Glu Lys Met Lys Ile Pro Phe Thr			
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Asp Ile Tyr Ile Gly Gly Ala Pro Pro Glu Ile Leu Gln Ser Arg Ala			
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Leu Arg Ala His Leu Pro Leu Asp Ile Asn Phe Arg Gly Cys Met Lys			
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Gly Phe Gln Phe Gln Lys Lys Asp Phe Asn Leu Leu Glu Gln Thr Glu			
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Thr Leu Gly Val Gly Tyr Gly Cys Pro Glu Asp Ser Leu Ile Ser Arg			
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Arg Ala Tyr Phe Asn Gly Gln Ser Phe Ile Ala Ser Ile Gln Lys Ile			
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Ser Phe Phe Asp Gly Phe Glu Gly Gly Phe Asn Phe Arg Thr Leu Gln	
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cca aat ggg tta cta ttc tat tat gct tca ggg tca gac gtg ttc tcc	3844
Pro Asn Gly Leu Leu Phe Tyr Tyr Ala Ser Gly Ser Asp Val Phe Ser	
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Ile Ser Leu Asp Asn Gly Thr Val Ile Met Asp Val Lys Gly Ile Lys	
1280 1285 1290	
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Val Gln Ser Val Asp Lys Gln Tyr Asn Asp Gly Leu Ser His Phe Val	
1295 1300 1305	
att agc tct gtc tca ccc aca aga tat gaa ctg ata gta gat aaa agc	3988
Ile Ser Ser Val Ser Pro Thr Arg Tyr Glu Leu Ile Val Asp Lys Ser	
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Arg Val Gly Ser Lys Asn Pro Thr Lys Gly Lys Ile Glu Gln Thr Gln	
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His Thr Ser Leu Tyr Glu Cys Pro Ile Glu Ser Ser Pro Leu Phe Leu	
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Ser Asn Ser Pro Arg Ala Ile Glu His Ala Tyr Gln Tyr Gly Gly Thr	
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Lys Ser Gln Phe Ser Ile Arg Leu Arg Thr Arg Ser Ser His Gly Met	
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Ser Thr Glu Gly Gly Tyr Val Val Leu Asp Glu Ser Phe Asn Ile Gly	
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Leu Lys Phe Glu Ile Ala Phe Glu Val Arg Pro Arg Ser Ser Ser Gly	
1665 1670 1675	
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Thr Leu Val His Gly His Ser Val Asn Gly Glu Tyr Leu Asn Val His	
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Met Lys Asn Gly Gln Val Ile Val Lys Val Asn Asn Gly Ile Arg Asp	
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His Arg Ile Thr Val Ile Arg Asp Ser Asn Val Val Gln Leu Asp Val
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 Asp Asp Asp Lys

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<212> PRT

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<400> 6

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Trp Ser Ala Ala Cys Ser Arg Ala Ala Ser Gly Asp Asp Asn Ala Phe
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Glu Thr Ser Glu Pro Arg Val Ala Leu Gly Arg Leu Pro Pro Ala Ala
 50 55 60

Glu Lys Cys Asn Ala Gly Phe Phe His Thr Leu Ser Gly Glu Cys Val
 65 70 75 80

Pro Cys Asp Cys Asn Gly Asn Ser Asn Glu Cys Leu Asp Gly Ser Gly
 85 90 95

Tyr Cys Val His Cys Gln Arg Asn Thr Thr Gly Glu His Cys Glu Lys
 100 105 110

Cys Leu Asp Gly Tyr Ile Gly Asp Ser Ile Arg Gly Ala Pro Gln Phe
 115 120 125

Cys Gln Pro Cys Pro Cys Pro Leu Pro His Leu Ala Asn Phe Pro Glu
 130 135 140

Ser Cys Tyr Arg Lys Asn Gly Ala Val Arg Cys Ile Cys Asn Glu Asn

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Pro	Phe	Leu	Ile	Gly	Ser	Thr	Cys	Lys	Lys	Cys	Asp	Cys	Ser	Gly	Asn
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Ser	Asp	Pro	Asn	Leu	Ile	Phe	Glu	Asp	Cys	Asp	Glu	Val	Thr	Gly	Gln
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Glu	Glu	Gly	Phe	Glu	Pro	Pro	Thr	Gly	Cys	Asp	Lys	Cys	Val	Trp	Asp
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Leu	Thr	Asp	Asp	Leu	Arg	Leu	Ala	Ala	Leu	Ser	Ile	Glu	Glu	Gly	Lys
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Gln	Lys	Met	Leu	Glu	Glu	Ile	Arg	Ser	Arg	Gln	Pro	Phe	Phe	Thr	Gln
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Arg	Glu	Leu	Val	Asp	Glu	Glu	Ala	Asp	Glu	Ala	Tyr	Glu	Leu	Leu	Ser
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Gln	Ala	Glu	Ser	Trp	Gln	Arg	Leu	His	Asn	Glu	Thr	Arg	Thr	Leu	Phe
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Pro	Val	Val	Leu	Glu	Gln	Leu	Asp	Asp	Tyr	Asn	Ala	Lys	Leu	Ser	Asp
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 Gln Glu Arg Val Arg Glu Gln Met Glu Val Val Asn Met Ser Leu Ser
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 Ala Lys Ser Glu Leu Gln Val Lys Leu Ser Asn Leu Ser Asn Leu Ser
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 His Asp Leu Val Gln Glu Ala Ile Asp His Ala Gln Asp Leu Gln Gln
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 Glu Ala Asn Glu Leu Ser Arg Lys Leu His Ser Ser Asp Met Asn Gly
 595 600 605
 Leu Val Gln Lys Ala Leu Asp Ala Ser Asn Val Tyr Glu Asn Ile Val
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 Asn Tyr Val Ser Glu Ala Asn Glu Thr Ala Glu Phe Ala Leu Asn Thr
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 Thr Asp Arg Ile Tyr Asp Ala Val Ser Gly Ile Asp Thr Gln Ile Ile
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 Tyr His Lys Asp Glu Ser Glu Asn Leu Leu Asn Gln Ala Arg Glu Leu
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 Gln Ala Lys Ala Glu Ser Ser Ser Asp Glu Ala Val Ala Asp Thr Ser
 675 680 685
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 690 695 700
 Leu Ser Asp Ala Val Lys Gln Leu Gln Ala Ala Glu Arg Gly Asp Ala
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 Thr Ala Val Asn Ser Ala Arg Asp Ala Val Arg Asn Leu Thr Glu Val
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 Val Pro Gln Leu Leu Asp Gln Leu Arg Thr Val Glu Gln Lys Arg Pro
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 Gln Thr Arg Ser Val Ala Ser Lys Ile Gln Val Ser Met Met Phe Asp
 820 825 830
 Gly Gln Ser Ala Val Glu Val His Ser Arg Thr Ser Met Asp Asp Leu
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 Lys Ala Phe Thr Ser Leu Ser Leu Tyr Met Lys Pro Pro Val Lys Arg
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His Asn Asp Lys Lys Met Ile Leu Val Val Asp Arg Arg His Val Lys 1140	1145	1150
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Leu Pro Leu Asp Ile Asn Phe Arg Gly Cys Met Lys Gly Phe Gln Phe 185	1190	1195
Gln Lys Lys Asp Phe Asn Leu Leu Glu Gln Thr Glu Thr Leu Gly Val 1205	1210	1215
Gly Tyr Gly Cys Pro Glu Asp Ser Leu Ile Ser Arg Arg Ala Tyr Phe 1220	1225	1230
Asn Gly Gln Ser Phe Ile Ala Ser Ile Gln Lys Ile Ser Phe Phe Asp 1235	1240	1245
Gly Phe Glu Gly Gly Phe Asn Phe Arg Thr Leu Gln Pro Asn Gly Leu 1250	1255	1260
Leu Phe Tyr Tyr Ala Ser Gly Ser Asp Val Phe Ser Ile Ser Leu Asp 265	1270	1275
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Asp Lys Gln Tyr Asn Asp Gly Leu Ser His Phe Val Ile Ser Ser Val 1300	1305	1310
Ser Pro Thr Arg Tyr Glu Leu Ile Val Asp Lys Ser Arg Val Gly Ser 1315	1320	1325
Lys Asn Pro Thr Lys Gly Lys Ile Glu Gln Thr Gln Ala Ser Glu Lys 1330	1335	1340
Lys Phe Tyr Phe Gly Gly Ser Pro Ile Ser Ala Gln Tyr Ala Asn Phe 345	1350	1355
Thr Gly Cys Ile Ser Asn Ala Tyr Phe Thr Arg Val Asp Arg Asp Val 1365	1370	1375
Glu Val Glu Asp Phe Gln Arg Tyr Thr Glu Lys Val His Thr Ser Leu 1380	1385	1390
Tyr Glu Cys Pro Ile Glu Ser Ser Pro Leu Phe Leu Leu His Lys Lys 1395	1400	1405
Gly Lys Asn Leu Ser Lys Pro Lys Ala Ser Gln Asn Lys Lys Gly Gly 1410	1415	1420
Lys Ser Lys Asp Ala Pro Ser Trp Asp Pro Val Ala Leu Lys Leu Pro 425	1430	1435
Glu Arg Asn Thr Pro Arg Asn Ser His Cys His Leu Ser Asn Ser Pro 1445	1450	1455

Arg Ala Ile Glu His Ala Tyr Gln Tyr Gly Gly Thr Ala Asn Ser Arg
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 Gln Glu Phe Glu His Leu Lys Gly Asp Phe Gly Ala Lys Ser Gln Phe
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 Ser Ile Arg Leu Arg Thr Arg Ser Ser His Gly Met Ile Phe Tyr Val
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 Gly His Ser Val Asn Gly Glu Tyr Leu Asn Val His Met Lys Asn Gly
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 Gln Val Ile Val Lys Val Asn Asn Gly Ile Arg Asp Phe Ser Thr Ser
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 Val Ile Arg Asp Ser Asn Val Val Gln Leu Asp Val Asp Ser Glu Val
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 Pro Val Phe Val Gly Gly Val Pro Glu Ser Leu Leu Thr Pro Arg Leu
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gcg gtt ggc agg caa gac ccg cct gag acg agc gaa ccc cgc gtg gct 96
 Ala Val Gly Arg Gln Asp Pro Pro Glu Thr Ser Glu Pro Arg Val Ala
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ctg gga cgc ctg ccg cct gcg gcc gag aaa tgc aat gct gga ttc ttt 144
 Leu Gly Arg Leu Pro Pro Ala Ala Glu Lys Cys Asn Ala Gly Phe Phe
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cac acc ctg tcg gga gaa tgt gtg ccc tgc gac tgt aat ggc aat tcc 192
 His Thr Leu Ser Gly Glu Cys Val Pro Cys Asp Cys Asn Gly Asn Ser
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 Asn Glu Cys Leu Asp Gly Ser Gly Tyr Cys Val His Cys Gln Arg Asn
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 Thr Thr Gly Glu His Cys Glu Lys Cys Leu Asp Gly Tyr Ile Gly Asp
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tcc atc agg gga gca ccc caa ttc tgc cag ccg tgc ccc tgt ccc ctg 336
 Ser Ile Arg Gly Ala Pro Gln Phe Cys Gln Pro Cys Pro Cys Pro Leu
 100 105 110

ccc cac ttg gcc aat ttt cca gaa tcc tgc tat agg aaa aat gga gct 384
 Pro His Leu Ala Asn Phe Pro Glu Ser Cys Tyr Arg Lys Asn Gly Ala
 115 120 125

gtt cgg tgc att tgt aac gaa aat tat gct gga cct aac tgt gaa aga 432
 Val Arg Cys Ile Cys Asn Glu Asn Tyr Ala Gly Pro Asn Cys Glu Arg
 130 135 140

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Cys 145	Ala	Pro	Gly	Tyr	Tyr 150	Gly	Asn	Pro	Phe	Leu 155	Ile	Gly	Ser	Thr	Cys 160	
aag	aaa	tgt	gac	tgc	agt	gga	aat	tca	gat	ccc	aac	ctg	atc	ttt	gaa	528
Lys	Lys	Cys	Asp	Cys	Ser	Gly	Asn	Ser	Asp	Pro	Asn	Leu	Ile	Phe	Glu	
				165					170					175		
gat	tgt	gat	gaa	gtc	act	ggc	cag	tgt	agg	aat	tgc	tta	cgc	aac	acc	576
Asp	Cys	Asp	Glu	Val	Thr	Gly	Gln	Cys	Arg	Asn	Cys	Leu	Arg	Asn	Thr	
			180					185					190			
acc	gga	ttc	aag	tgt	gaa	cgt	tgc	gct	cct	ggc	tac	tat	ggg	gac	gcc	624
Thr	Gly	Phe	Lys	Cys	Glu	Arg	Cys	Ala	Pro	Gly	Tyr	Tyr	Gly	Asp	Ala	
		195					200					205				
agg	ata	gcc	aag	aac	tgt	gca	gtg	tgc	aac	tgc	ggg	gga	ggc	cca	tgt	672
Arg	Ile	Ala	Lys	Asn	Cys	Ala	Val	Cys	Asn	Cys	Gly	Gly	Gly	Pro	Cys	
	210					215					220					
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Asp	Ser	Val	Thr	Gly	Glu	Cys	Leu	Glu	Glu	Gly	Phe	Glu	Pro	Pro	Thr	
225					230					235					240	
ggc	tgt	gat	aag	tgc	gtc	tgg	gac	ctg	act	gat	gac	ctg	cgg	tta	gca	768
Gly	Cys	Asp	Lys	Cys	Val	Trp	Asp	Leu	Thr	Asp	Asp	Leu	Arg	Leu	Ala	
				245				250						255		
gcg	ctc	tcc	atc	gag	gaa	ggc	aaa	tcc	ggg	gtg	ctg	agc	gta	tcc	tct	816
Ala	Leu	Ser	Ile	Glu	Glu	Gly	Lys	Ser	Gly	Val	Leu	Ser	Val	Ser	Ser	
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ggg	gcc	gcc	gct	cat	agg	cac	gtg	aat	gaa	atc	aac	gcc	acc	atc	tac	864
Gly	Ala	Ala	Ala	His	Arg	His	Val	Asn	Glu	Ile	Asn	Ala	Thr	Ile	Tyr	
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ctc	ctc	aaa	aca	aaa	ttg	tca	gaa	aga	gaa	aac	caa	tac	gcc	cta	aga	912
Leu	Leu	Lys	Thr	Lys	Leu	Ser	Glu	Arg	Glu	Asn	Gln	Tyr	Ala	Leu	Arg	
	290					295					300					
aag	ata	caa	atc	aac	aat	gct	gag	aac	acg	atg	aaa	agc	ctt	ctg	tct	960
Lys	Ile	Gln	Ile	Asn	Asn	Ala	Glu	Asn	Thr	Met	Lys	Ser	Leu	Leu	Ser	
305					310					315				320		
gac	gta	gag	gaa	tta	gtt	gaa	aag	gaa	aat	caa	gcc	tcc	aga	aaa	gga	1008
Asp	Val	Glu	Glu	Leu	Val	Glu	Lys	Glu	Asn	Gln	Ala	Ser	Arg	Lys	Gly	
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caa	ctt	gtt	cag	aag	gaa	agc	atg	gac	acc	att	aac	cac	gca	agt	cag	1056
Gln	Leu	Val	Gln	Lys	Glu	Ser	Met	Asp	Thr	Ile	Asn	His	Ala	Ser	Gln	
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ctg	gta	gag	caa	gcc	cat	gat	atg	agg	gat	aaa	atc	caa	gag	atc	aac	1104
Leu	Val	Glu	Gln	Ala	His	Asp	Met	Arg	Asp	Lys	Ile	Gln	Glu	Ile	Asn	
		355					360					365				
aac	aag	atg	ctc	tat	tat	ggg	gaa	gag	cat	gaa	ctt	agc	ccc	aag	gaa	1152
Asn	Lys	Met	Leu	Tyr	Tyr	Gly	Glu	Glu	His	Glu	Leu	Ser	Pro	Lys	Glu	
	370					375					380					
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Ile	Ser	Glu	Lys	Leu	Val	Leu	Ala	Gln	Lys	Met	Leu	Glu	Glu	Ile	Arg	

385		390		395		400	
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Ser Arg Gln Pro Phe Phe Thr Gln Arg Glu Leu Val Asp Glu Glu Ala							
		405		410		415	
gat gag gct tac gaa cta ctg agc cag gct gag agc tgg cag cgg ctg	1296						
Asp Glu Ala Tyr Glu Leu Leu Ser Gln Ala Glu Ser Trp Gln Arg Leu							
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cac aat gag acc cgc act ctg ttt cct gtc gtc ctg gag cag ctg gat	1344						
His Asn Glu Thr Arg Thr Leu Phe Pro Val Val Leu Glu Gln Leu Asp							
		435		440		445	
gac tac aat gct aag ttg tca gat ctc cag gaa gca ctt gac cag gcc	1392						
Asp Tyr Asn Ala Lys Leu Ser Asp Leu Gln Glu Ala Leu Asp Gln Ala							
		450		455		460	
ctt aac tat gtc agg gat gcc gaa gac atg aac agg gcc aca gca gcc	1440						
Leu Asn Tyr Val Arg Asp Ala Glu Asp Met Asn Arg Ala Thr Ala Ala							
		465		470		475	
agg cag cgg gac cat gag aaa caa cag gaa aga gtg agg gaa caa atg	1488						
Arg Gln Arg Asp His Glu Lys Gln Gln Glu Arg Val Arg Glu Gln Met							
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gaa gtg gtg aac atg tct ctg agc aca tct gcg gac tct ctg aca aca	1536						
Glu Val Val Asn Met Ser Leu Ser Thr Ser Ala Asp Ser Leu Thr Thr							
		500		505		510	
cct cgt cta act ctt tca gaa ctt gat gat ata ata aag aat gcg tca	1584						
Pro Arg Leu Thr Leu Ser Glu Leu Asp Asp Ile Ile Lys Asn Ala Ser							
		515		520		525	
ggg att tat gca gaa ata gat gga gcc aaa agt gaa cta caa gta aaa	1632						
Gly Ile Tyr Ala Glu Ile Asp Gly Ala Lys Ser Glu Leu Gln Val Lys							
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cta tct aac cta agt aac ctc agc cat gat tta gtc caa gaa gct att	1680						
Leu Ser Asn Leu Ser Asn Leu Ser His Asp Leu Val Gln Glu Ala Ile							
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gac cat gca cag gac ctt caa caa gaa gct aat gaa ttg agc agg aag	1728						
Asp His Ala Gln Asp Leu Gln Gln Glu Ala Asn Glu Leu Ser Arg Lys							
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ttg cac agt tca gat atg aac ggg ctg gta cag aag gct ttg gat gca	1776						
Leu His Ser Ser Asp Met Asn Gly Leu Val Gln Lys Ala Leu Asp Ala							
		580		585		590	
tca aat gtc tat gaa aat att gtt aat tat gtt agt gaa gcc aat gaa	1824						
Ser Asn Val Tyr Glu Asn Ile Val Asn Tyr Val Ser Glu Ala Asn Glu							
		595		600		605	
aca gca gaa ttt gct ttg aac acc act gac cga att tat gat gcg gtg	1872						
Thr Ala Glu Phe Ala Leu Asn Thr Thr Asp Arg Ile Tyr Asp Ala Val							
		610		615		620	
agt ggg att gat act caa atc att tac cat aaa gat gaa agt gag aac	1920						
Ser Gly Ile Asp Thr Gln Ile Ile Tyr His Lys Asp Glu Ser Glu Asn							
		625		630		635	
						640	

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Asp Glu Ala Val Ala Asp Thr Ser Arg Arg Val Gly Gly Ala Leu Ala	
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agg aaa agt gcc ctt aaa acc aga ctc agt gat gcc gtt aag caa cta	2064
Arg Lys Ser Ala Leu Lys Thr Arg Leu Ser Asp Ala Val Lys Gln Leu	
675 680 685	
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Gln Ala Ala Glu Arg Gly Asp Ala Gln Gln Arg Leu Gly Gln Ser Arg	
690 695 700	
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Leu Ile Thr Glu Glu Ala Asn Arg Thr Thr Met Glu Val Gln Gln Ala	
705 710 715 720	
act gcc ccc atg gcc aac aat cta acc aac tgg tca cag aat ctt caa	2208
Thr Ala Pro Met Ala Asn Asn Leu Thr Asn Trp Ser Gln Asn Leu Gln	
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His Phe Asp Ser Ser Ala Tyr Asn Thr Ala Val Asn Ser Ala Arg Asp	
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Ala Val Arg Asn Leu Thr Glu Val Val Pro Gln Leu Leu Asp Gln Leu	
755 760 765	
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Arg Thr Val Glu Gln Lys Arg Pro Ala Ser Asn Val Ser Ala Ser Ile	
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Gln Arg Ile Arg Glu Leu Ile Ala Gln Thr Arg Ser Val Ala Ser Lys	
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Ser Arg Thr Ser Met Asp Asp Leu Lys Ala Phe Thr Ser Leu Ser Leu	
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Tyr Met Lys Pro Pro Val Lys Arg Pro Glu Leu Thr Glu Thr Ala Asp	
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Gly Leu Ala Ile Lys Asn Asp Asn Leu Val Tyr Val Tyr Asn Leu Gly	
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Thr Lys Asp Val Glu Ile Pro Leu Asp Ser Lys Pro Val Ser Ser Trp	
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Pro Ala Tyr Phe Ser Ile Val Lys Ile Glu Arg Val Gly Lys His Gly	
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Lys Val Phe Leu Thr Val Pro Ser Leu Ser Ser Thr Ala Glu Glu Lys	
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Asn Gly Tyr Leu His Val Phe Tyr Asp Phe Gly Phe Ser Ser Gly Arg	
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Val His Leu Glu Asp Thr Leu Lys Lys Ala Gln Ile Asn Asp Ala Lys	
1090 1095 1100	
tac cat gag atc tca atc att tac cac aat gat aag aaa atg atc ttg	3360
Tyr His Glu Ile Ser Ile Ile Tyr His Asn Asp Lys Lys Met Ile Leu	
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Val Val Asp Arg Arg His Val Lys Ser Met Asp Asn Glu Lys Met Lys	
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Ile Pro Phe Thr Asp Ile Tyr Ile Gly Gly Ala Pro Pro Glu Ile Leu	
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Lys Gly Ile Lys Val Gln Ser Val Asp Lys Gln Tyr Asn Asp Gly Leu	
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Ser His Phe Val Ile Ser Ser Val Ser Pro Thr Arg Tyr Glu Leu Ile	
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Ala Ser Gln Asn Lys Lys Gly Gly Lys Ser Lys Asp Ala Pro Ser Trp			
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His Cys His Leu Ser Asn Ser Pro Arg Ala Ile Glu His Ala Tyr Gln			
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Tyr Gly Gly Thr Ala Asn Ser Arg Gln Glu Phe Glu His Leu Lys Gly			
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Asp Phe Gly Ala Lys Ser Gln Phe Ser Ile Arg Leu Arg Thr Arg Ser			
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Ser His Gly Met Ile Phe Tyr Val Ser Asp Gln Glu Glu Asn Asp Phe			
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Met Thr Leu Phe Leu Ala His Gly Arg Leu Val Tyr Met Phe Asn Val			
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Val Ile Asp Gly Leu Arg Val Leu Glu Glu Ser Leu Pro Pro Thr Glu			
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Gly Lys Ala Val Lys Asn Val Gln Ile Asn Ser Ile Tyr Ser Phe Ser			
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Gly Cys Leu Ser Asn Leu Gln Leu Asn Gly Ala Ser Ile Thr Ser Ala			
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tct cag aca ttc agt gtg acc cct tgc ttt gaa ggc ccc atg gaa aca			4848
Ser Gln Thr Phe Ser Val Thr Pro Cys Phe Glu Gly Pro Met Glu Thr			
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 Gly Thr Tyr Phe Ser Thr Glu Gly Gly Tyr Val Val Leu Asp Glu Ser
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ttc aat att gga ttg aag ttt gaa att gca ttt gaa gtc cgt ccc aga 4944
 Phe Asn Ile Gly Leu Lys Phe Glu Ile Ala Phe Glu Val Arg Pro Arg
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agc agt tcc gga acc ctg gtc cac ggc cac agt gtc aat ggg gag tac 4992
 Ser Ser Ser Gly Thr Leu Val His Gly His Ser Val Asn Gly Glu Tyr
 1650 1655 1660

cta aat gtt cac atg aaa aat gga cag gtc ata gtg aaa gtc aat aat 5040
 Leu Asn Val His Met Lys Asn Gly Gln Val Ile Val Lys Val Asn Asn
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 Gly Ile Arg Asp Phe Ser Thr Ser Val Thr Pro Lys Gln Ser Leu Cys
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gat ggc aga tgg cac aga att aca gtt att aga gat tct aat gtg gtt 5136
 Asp Gly Arg Trp His Arg Ile Thr Val Ile Arg Asp Ser Asn Val Val
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 Gln Leu Asp Val Asp Ser Glu Val Asn His Val Val Gly Pro Leu Asn
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gaa tct cta ctg aca cca cgc ttg gcc ccc agc aaa ccc ttc aca ggc 5280
 Glu Ser Leu Leu Thr Pro Arg Leu Ala Pro Ser Lys Pro Phe Thr Gly
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 Cys Ile Arg His Phe Val Ile Asp Gly His Pro Val Ser Phe Ser Lys
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gca gcc ctg gtc agc ggc gcc gta agc atc aac tcc tgt cca gca gcc 5376
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 His Thr Leu Ser Gly Glu Cys Val Pro Cys Asp Cys Asn Gly Asn Ser
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 Thr Thr Gly Glu His Cys Glu Lys Cys Leu Asp Gly Tyr Ile Gly Asp
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 Ser Ile Arg Gly Ala Pro Gln Phe Cys Gln Pro Cys Pro Cys Pro Leu
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 Pro His Leu Ala Asn Phe Pro Glu Ser Cys Tyr Arg Lys Asn Gly Ala
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 Val Arg Cys Ile Cys Asn Glu Asn Tyr Ala Gly Pro Asn Cys Glu Arg
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 Gly Cys Asp Lys Cys Val Trp Asp Leu Thr Asp Asp Leu Arg Leu Ala
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 Gly Ala Ala Ala His Arg His Val Asn Glu Ile Asn Ala Thr Ile Tyr
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 Asp Val Glu Glu Leu Val Glu Lys Glu Asn Gln Ala Ser Arg Lys Gly
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 Gln Leu Val Gln Lys Glu Ser Met Asp Thr Ile Asn His Ala Ser Gln
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Leu Val Glu Gln Ala His Asp Met Arg Asp Lys Ile Gln Glu Ile Asn
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 Asn Lys Met Leu Tyr Tyr Gly Glu Glu His Glu Leu Ser Pro Lys Glu
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 Ile Ser Glu Lys Leu Val Leu Ala Gln Lys Met Leu Glu Glu Ile Arg
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 Thr Ala Glu Phe Ala Leu Asn Thr Thr Asp Arg Ile Tyr Asp Ala Val
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 Ser Gly Ile Asp Thr Gln Ile Ile Tyr His Lys Asp Glu Ser Glu Asn
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 Arg Lys Ser Ala Leu Lys Thr Arg Leu Ser Asp Ala Val Lys Gln Leu

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Thr Ala Pro Met Ala Asn Asn Leu Thr Asn Trp Ser Gln Asn Leu Gln		
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Arg Thr Val Glu Gln Lys Arg Pro Ala Ser Asn Val Ser Ala Ser Ile		
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Gln Arg Ile Arg Glu Leu Ile Ala Gln Thr Arg Ser Val Ala Ser Lys		
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Ile Gln Val Ser Met Met Phe Asp Gly Gln Ser Ala Val Glu Val His		
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Ser Arg Thr Ser Met Asp Asp Leu Lys Ala Phe Thr Ser Leu Ser Leu		
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Tyr Met Lys Pro Pro Val Lys Arg Pro Glu Leu Thr Glu Thr Ala Asp		
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Gln Phe Ile Leu Tyr Leu Gly Ser Lys Asn Ala Lys Lys Glu Tyr Met		
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Gly Leu Ala Ile Lys Asn Asp Asn Leu Val Tyr Val Tyr Asn Leu Gly		
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Thr Lys Asp Val Glu Ile Pro Leu Asp Ser Lys Pro Val Ser Ser Trp		
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Pro Ala Tyr Phe Ser Ile Val Lys Ile Glu Arg Val Gly Lys His Gly		
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Lys Val Phe Leu Thr Val Pro Ser Leu Ser Ser Thr Ala Glu Glu Lys		
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Phe Ile Lys Lys Gly Glu Phe Ser Gly Asp Asp Ser Leu Leu Asp Leu		
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Asp Pro Glu Asp Thr Val Phe Tyr Val Gly Gly Val Pro Ser Asn Phe		
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Ile Tyr Asn Met Asp Pro Ser Thr Ser Val Pro Cys Ala Arg Asp Lys		
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Asp Gly Arg Trp His Arg Ile Thr Val Ile Arg Asp Ser Asn Val Val		
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Ile Leu Val Val Asp Arg Arg His Val Lys Ser Thr Asp Asn Glu Lys	
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Lys Lys Ile Pro Phe Thr Asp Ile Tyr Ile Gly Gly Ala Pro Gln Glu	
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Ala Ser Ile Gln Lys Ile Ser Phe Phe Asp Gly Phe Glu Gly Gly Phe	
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Leu Val Val Asp Lys Ser Arg Leu Arg Gly Lys Asn Pro Thr Lys Gly	
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Arg Tyr Ser Glu Lys Val His Thr Ser Leu Tyr Glu Cys Pro Ile Glu	
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Cys Ile Arg His Phe Val Ile Asp Ser Arg Pro Val Ser Phe Ser Lys			
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Gly Ile Tyr Ala Glu Ile Asp Gly Ala Lys Asn Glu Leu Gln Gly Lys	
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cta tcc aac ctg agt aac ctc agt cat gac ttg gtt cag gaa gct acg	1680
Leu Ser Asn Leu Ser Asn Leu Ser His Asp Leu Val Gln Glu Ala Thr	
545 550 555 560	
gac cat gca tac aat ctt caa cag gaa gcc gat gag cta agc aga aat	1728
Asp His Ala Tyr Asn Leu Gln Gln Glu Ala Asp Glu Leu Ser Arg Asn	
565 570 575	
ttg cac agt tca gac atg aac ggg ctg gta cag aag gct ttg gat gca	1776
Leu His Ser Ser Asp Met Asn Gly Leu Val Gln Lys Ala Leu Asp Ala	
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tca aac gtc tat gaa aat atc gcc aat tat gtc agt gaa gcc aac gaa	1824
Ser Asn Val Tyr Glu Asn Ile Ala Asn Tyr Val Ser Glu Ala Asn Glu	
595 600 605	
aca gca gaa ctt gct ctg aat atc act gat cga att tat gat gct gtg	1872
Thr Ala Glu Leu Ala Leu Asn Ile Thr Asp Arg Ile Tyr Asp Ala Val	
610 615 620	
agt ggg att gac acg cag atc att tac cat aag gat gaa agt gac aac	1920
Ser Gly Ile Asp Thr Gln Ile Ile Tyr His Lys Asp Glu Ser Asp Asn	
625 630 635 640	
ctt ctc aat caa gcc aga gag ctg cag gcc aag gca gat tca tgc aat	1968
Leu Leu Asn Gln Ala Arg Glu Leu Gln Ala Lys Ala Asp Ser Cys Asn	
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gat gaa gca gtg gct gac acc agc agg cgt gtg ggt gga gcc ctg tgg	2016

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Arg	Lys	Gly	Ala	Leu	Arg	Asp	Arg	Leu	Asn	Asp	Ala	Val	Lys	Gln	Leu	
		675					680					685				
cag	gca	gca	gag	aga	ggg	gac	gcc	cac	cag	cgc	ctg	ggc	cag	tcc	aag	2112
Gln	Ala	Ala	Glu	Arg	Gly	Asp	Ala	His	Gln	Arg	Leu	Gly	Gln	Ser	Lys	
	690					695					700					
ctc	ttc	att	gag	gaa	gct	aac	aag	acg	aca	gcg	gct	gtc	caa	cag	gtt	2160
Leu	Phe	Ile	Glu	Glu	Ala	Asn	Lys	Thr	Thr	Ala	Ala	Val	Gln	Gln	Val	
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acc	aca	cca	atg	gct	aac	aac	ctc	agc	aac	tgg	tcc	cag	aac	ctg	cag	2208
Thr	Thr	Pro	Met	Ala	Asn	Asn	Leu	Ser	Asn	Trp	Ser	Gln	Asn	Leu	Gln	
			725					730					735			
acc	ttt	gac	tca	tct	gca	tat	aac	act	gca	gtg	gac	tct	gct	cgg	gac	2256
Thr	Phe	Asp	Ser	Ser	Ala	Tyr	Asn	Thr	Ala	Val	Asp	Ser	Ala	Arg	Asp	
			740				745						750			
gca	gtg	aga	aac	ctc	acc	gag	gtt	gtc	ccc	cag	ctt	ctg	gat	cag	ctt	2304
Ala	Val	Arg	Asn	Leu	Thr	Glu	Val	Val	Pro	Gln	Leu	Leu	Asp	Gln	Leu	
		755				760						765				
cgt	act	gtg	gag	cag	aag	cgg	cct	gca	agc	aac	att	tct	gcc	agc	atc	2352
Arg	Thr	Val	Glu	Gln	Lys	Arg	Pro	Ala	Ser	Asn	Ile	Ser	Ala	Ser	Ile	
	770					775					780					
cag	agc	atc	cga	gag	ctc	att	gct	caa	acc	agg	agt	gtc	gca	agc	aag	2400
Gln	Ser	Ile	Arg	Glu	Leu	Ile	Ala	Gln	Thr	Arg	Ser	Val	Ala	Ser	Lys	
785				790					795						800	
atc	caa	gtc	tcc	atg	atg	ttt	gat	ggc	cag	tca	gct	gtc	gaa	gtg	cac	2448
Ile	Gln	Val	Ser	Met	Met	Phe	Asp	Gly	Gln	Ser	Ala	Val	Glu	Val	His	
			805					810					815			
ccc	aaa	gtc	agt	gtg	gat	gac	ctg	aag	gcc	ttc	aca	tcc	atc	agc	ttg	2496
Pro	Lys	Val	Ser	Val	Asp	Asp	Leu	Lys	Ala	Phe	Thr	Ser	Ile	Ser	Leu	
			820				825						830			
tac	atg	aag	cct	cct	cca	aag	ccg	gca	gag	ccc	act	ggg	gcc	tgg	gta	2544
Tyr	Met	Lys	Pro	Pro	Pro	Lys	Pro	Ala	Glu	Pro	Thr	Gly	Ala	Trp	Val	
		835				840						845				
gca	gat	cag	ttt	gtc	ctc	tac	ctc	gga	agc	aaa	aac	gcc	aaa	aaa	gaa	2592
Ala	Asp	Gln	Phe	Val	Leu	Tyr	Leu	Gly	Ser	Lys	Asn	Ala	Lys	Lys	Glu	
	850					855					860					
tac	atg	ggg	ctg	gca	atc	aaa	aat	gat	aac	ctg	gta	tac	gtt	tac	aat	2640
Tyr	Met	Gly	Leu	Ala	Ile	Lys	Asn	Asp	Asn	Leu	Val	Tyr	Val	Tyr	Asn	
865				870						875					880	
ttg	ggg	atg	aaa	gat	gtg	gaa	att	ctc	ctg	gat	tcc	aag	cct	gtg	agc	2688
Leu	Gly	Met	Lys	Asp	Val	Glu	Ile	Leu	Leu	Asp	Ser	Lys	Pro	Val	Ser	
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tcc	tgg	ccc	gct	tac	ttt	agt	att	gtc	aag	att	gaa	agg	gta	ggg	gaa	2736
Ser	Trp	Pro	Ala	Tyr	Phe	Ser	Ile	Val	Lys	Ile	Glu	Arg	Val	Gly	Glu	

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His	Gly	Lys	Val	Phe	Leu	Thr	Val	Pro	Ser	Leu	Ser	Ser	Thr	Ala	Glu			
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gaa	aag	ttt	att	aag	aag	ggg	gag	ttt	gca	gga	gat	gac	tcc	ttg	ctg	2832		
Glu	Lys	Phe	Ile	Lys	Lys	Gly	Glu	Phe	Ala	Gly	Asp	Asp	Ser	Leu	Leu			
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gat	gtg	acc	cct	gag	gac	act	gtg	ttt	tac	gtt	ggg	ggg	gtg	cct	gcg	2880		
Asp	Val	Thr	Pro	Glu	Asp	Thr	Val	Phe	Tyr	Val	Gly	Gly	Val	Pro	Ala			
945						950						955						960
aac	ttc	aag	ctc	cct	gcc	agc	tta	aac	ctg	ccc	agc	tac	tca	ggc	tgc	2928		
Asn	Phe	Lys	Leu	Pro	Ala	Ser	Leu	Asn	Leu	Pro	Ser	Tyr	Ser	Gly	Cys			
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Leu	Glu	Leu	Ala	Thr	Leu	Asn	Asn	Asp	Val	Ile	Ser	Leu	Tyr	Asn	Phe			
980						985						990						
aag	cac	atc	tat	aat	atg	gat	cca	tca	aag	tca	gtg	ccc	tgt	gcc	agg	3024		
Lys	His	Ile	Tyr	Asn	Met	Asp	Pro	Ser	Lys	Ser	Val	Pro	Cys	Ala	Arg			
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Gly	Gln	Val	Thr	Arg	Phe	Asp	Ile	Glu	Ile	Arg	Thr	Pro	Ala	Asp	Asn			
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atg	cgc	aat	ggc	tac	cta	cat	gtg	ttc	tat	gac	ttt	gga	ttc	agc	aat	3264		
Met	Arg	Asn	Gly	Tyr	Leu	His	Val	Phe	Tyr	Asp	Phe	Gly	Phe	Ser	Asn			
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ggc	ccc	gtg	cat	ctt	gaa	gac	acg	ttg	aaa	aaa	gcc	cag	att	aat	gat	3312		
Gly	Pro	Val	His	Leu	Glu	Asp	Thr	Leu	Lys	Lys	Ala	Gln	Ile	Asn	Asp			
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gcg	aaa	tat	cat	gag	atc	tca	atc	att	tat	cac	aac	gac	aaa	aaa	atg	3360		
Ala	Lys	Tyr	His	Glu	Ile	Ser	Ile	Ile	Tyr	His	Asn	Asp	Lys	Lys	Met			
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att	ttg	gtg	gtg	gac	aga	cgg	cac	gtt	aag	agc	aca	gac	aat	gag	aag	3408		
Ile	Leu	Val	Val	Asp	Arg	Arg	His	Val	Lys	Ser	Thr	Asp	Asn	Glu	Lys			
1125						1130						1135						
aaa	aag	att	cct	ttc	acg	gac	atc	tac	atc	gga	ggt	gcg	ccc	caa	gaa	3456		
Lys	Lys	Ile	Pro	Phe	Thr	Asp	Ile	Tyr	Ile	Gly	Gly	Ala	Pro	Gln	Glu			
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Val Leu Gln Ser Arg Thr Leu Arg Ala His Leu Pro Leu Asp Ile Asn	
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Phe Arg Gly Cys Met Lys Gly Phe Gln Phe Gln Lys Lys Asp Phe Asn	
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gct tca att cag aaa ata tct ttc ttt gat ggc ttt gaa gga ggc ttc	3696
Ala Ser Ile Gln Lys Ile Ser Phe Phe Asp Gly Phe Glu Gly Gly Phe	
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Asn Phe Arg Thr Leu Gln Pro Asn Gly Leu Leu Phe Tyr Tyr Thr Ser	
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Gly Ser Asp Val Phe Ser Ile Ser Leu Asp Asn Gly Thr Val Val Met	
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Asp Val Lys Gly Ile Lys Val Met Ser Thr Asp Lys Gln Tyr His Asp	
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Gly Leu Pro His Phe Val Val Thr Ser Ile Ser Asp Thr Arg Tyr Glu	
1285 1290 1295	
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Leu Val Val Asp Lys Ser Arg Leu Arg Gly Lys Asn Pro Thr Lys Gly	
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Lys Ala Glu Gln Thr Gln Thr Thr Glu Lys Lys Phe Tyr Phe Gly Gly	
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Ser Pro Ile Ser Pro Gln Tyr Ala Asn Phe Thr Gly Cys Ile Ser Asn	
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gcc tac ttt acc agg ttg gat aga gat gtg gaa gtc gaa gac ttc cag	4080
Ala Tyr Phe Thr Arg Leu Asp Arg Asp Val Glu Val Glu Asp Phe Gln	
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cgc tat tct gaa aag gtc cac act tca ctc tat gag tgt ccg att gag	4128
Arg Tyr Ser Glu Lys Val His Thr Ser Leu Tyr Glu Cys Pro Ile Glu	
1365 1370 1375	
tcg tca cct ctg ttt ctc ctt cac aaa aaa gga aag aat tcc tca aag	4176
Ser Ser Pro Leu Phe Leu Leu His Lys Lys Gly Lys Asn Ser Ser Lys	
1380 1385 1390	

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 Pro Lys Thr Asn Lys Gln Gly Glu Lys Ser Lys Asp Ala Pro Ser Trp
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 Asp Pro Ile Gly Leu Lys Phe Leu Glu Gln Lys Ala Pro Arg Asp Ser
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 His Cys His Leu Ser Ser Ser Pro Arg Ala Ile Glu His Ala Tyr Gln
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 Tyr Gly Gly Thr Ala Asn Ser Arg Gln Glu Phe Glu His Glu Gln Gly
 1445 1450 1455

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 Asp Phe Gly Glu Lys Ser Gln Phe Ala Ile Arg Leu Lys Thr Arg Ser
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 Ser His Gly Met Ile Phe Tyr Val Ser Asp Gln Glu Glu Asn Asp Phe
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 Met Thr Leu Phe Leu Ala His Gly Arg Leu Val Phe Met Phe Asn Val
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 Gly His Lys Lys Leu Lys Ile Arg Ser Gln Glu Lys Tyr Asn Asp Gly
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 Val Ile Asp Gly Leu Arg Val Leu Glu Glu Arg Leu Pro Pro Ser Gly
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 Ala Ala Trp Lys Ile Lys Gly Pro Ile Tyr Leu Gly Gly Val Ala Pro
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 Gly Thr Tyr Phe Ser Thr Glu Gly Gly Tyr Val Val Leu Asp Glu Ser
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 Leu Asn Val His Met Arg Asn Gly Gln Val Ile Val Lys Val Asn Asn
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 35 40 45

Arg Thr Leu Ser Gly Glu Cys Ala Pro Cys Asp Cys Asn Gly Asn Ser
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 His Glu Cys Leu Asp Gly Ser Gly Phe Cys Leu His Cys Gln Arg Asn
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 Thr Thr Gly Glu His Cys Glu Lys Cys Leu Asp Gly Tyr Ile Gly Asp
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 Ser Ile Arg Gly Thr Pro Arg Phe Cys Gln Pro Cys Pro Cys Pro Leu
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 Pro His Leu Ala Asn Phe Ala Glu Ser Cys Tyr Arg Lys Asn Gly Ala
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 Val Arg Cys Ile Cys Lys Glu Asn Tyr Val Gly Pro Asn Cys Glu Arg
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 Cys Ala Pro Gly Tyr Tyr Gly Asn Pro Leu Leu Ile Gly Ser Thr Cys
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 Lys Lys Cys Asp Cys Ser Gly Asn Ser Asp Pro Asn Leu Ile Phe Glu
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 Thr Gly Phe Lys Cys Glu Arg Cys Ala Pro Gly Tyr Tyr Gly Asp Ala
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 Arg Thr Ala Lys Asn Cys Ala Val Cys Asn Cys Gly Gly Gly Pro Arg
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 Asp Ser Val Thr Gly Glu Cys Leu Glu Glu Gly Phe Glu Val Pro Thr
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 Gly Cys Asp Lys Cys Val Trp Asp Leu Thr Asp Asp Leu Arg Leu Ala
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 Ala Leu Ser Ile Glu Glu Ser Lys Ser Gly Leu Leu Ser Val Ser Ser
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 Ala Ala Ala Ala His Arg His Val Thr Asp Met Asn Ser Thr Ile His
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 Leu Leu Arg Thr Arg Leu Ser Glu Arg Glu Asn Gln Tyr Thr Leu Arg
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 Lys Ile Gln Ile Asn Asn Ser Glu Asn Thr Leu Arg Ser Leu Leu Pro
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 Asp Val Glu Gly Leu His Glu Lys Gly Ser Gln Ala Ser Arg Lys Gly
 325 330 335
 Met Leu Val Glu Lys Glu Ser Met Asp Thr Ile Asp Gln Ala Thr His
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 Leu Val Glu Gln Ala His Asn Met Arg Asp Lys Ile Gln Glu Ile Asn
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 Ser Lys Met Leu Tyr Tyr Gly Glu Asn Gln Glu Leu Gly Pro Glu Glu

370		375		380
Ile Ala Glu Lys Leu Val Leu Ala Gln Lys Met Leu Glu Glu Ile Arg				
385		390		395
Ser Arg Gln Pro Phe Leu Thr His Arg Glu Leu Val Asp Glu Glu Ala				
	405		410	415
Asp Glu Ala Gln Glu Leu Leu Ser Gln Ala Glu Asn Trp Gln Arg Leu				
	420		425	430
His Asn Asp Thr Arg Ser Leu Phe Pro Val Val Leu Glu Gln Leu Asp				
	435		440	445
Asp Tyr Asn Ala Lys Leu Ser Asp Leu Gln Glu Ser Ile Asn Gln Ala				
	450		455	460
Leu Asp His Val Arg Asp Ala Glu Asp Met Asn Arg Ala Ile Thr Phe				
	465		470	475
Lys Gln Arg Asp His Glu Lys Gln His Glu Arg Val Lys Glu Gln Met				
	485		490	495
Glu Val Val Gly Ala Ser Leu Ser Met Ser Ala Asp Ser Leu Thr Ile				
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Pro Gln Leu Thr Leu Glu Glu Leu Asp Glu Ile Ile Lys Asn Ala Ser				
	515		520	525
Gly Ile Tyr Ala Glu Ile Asp Gly Ala Lys Asn Glu Leu Gln Gly Lys				
	530		535	540
Leu Ser Asn Leu Ser Asn Leu Ser His Asp Leu Val Gln Glu Ala Thr				
	545		550	555
Asp His Ala Tyr Asn Leu Gln Gln Glu Ala Asp Glu Leu Ser Arg Asn				
	565		570	575
Leu His Ser Ser Asp Met Asn Gly Leu Val Gln Lys Ala Leu Asp Ala				
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Ser Asn Val Tyr Glu Asn Ile Ala Asn Tyr Val Ser Glu Ala Asn Glu				
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Thr Ala Glu Leu Ala Leu Asn Ile Thr Asp Arg Ile Tyr Asp Ala Val				
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Ser Gly Ile Asp Thr Gln Ile Ile Tyr His Lys Asp Glu Ser Asp Asn				
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Leu Leu Asn Gln Ala Arg Glu Leu Gln Ala Lys Ala Asp Ser Cys Asn				
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Asp Glu Ala Val Ala Asp Thr Ser Arg Arg Val Gly Gly Ala Leu Trp				
	660		665	670
Arg Lys Gly Ala Leu Arg Asp Arg Leu Asn Asp Ala Val Lys Gln Leu				
	675		680	685
Gln Ala Ala Glu Arg Gly Asp Ala His Gln Arg Leu Gly Gln Ser Lys				
	690		695	700

Leu Phe Ile Glu Glu Ala Asn Lys Thr Thr Ala Ala Val Gln Gln Val
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 Thr Thr Pro Met Ala Asn Asn Leu Ser Asn Trp Ser Gln Asn Leu Gln
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 Thr Phe Asp Ser Ser Ala Tyr Asn Thr Ala Val Asp Ser Ala Arg Asp
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 770 775 780
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 785 790 795 800
 Ile Gln Val Ser Met Met Phe Asp Gly Gln Ser Ala Val Glu Val His
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 Pro Lys Val Ser Val Asp Asp Leu Lys Ala Phe Thr Ser Ile Ser Leu
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 Tyr Met Lys Pro Pro Pro Lys Pro Ala Glu Pro Thr Gly Ala Trp Val
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 Ala Asp Gln Phe Val Leu Tyr Leu Gly Ser Lys Asn Ala Lys Lys Glu
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 Tyr Met Gly Leu Ala Ile Lys Asn Asp Asn Leu Val Tyr Val Tyr Asn
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 Leu Gly Met Lys Asp Val Glu Ile Leu Leu Asp Ser Lys Pro Val Ser
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 Ser Trp Pro Ala Tyr Phe Ser Ile Val Lys Ile Glu Arg Val Gly Glu
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 His Gly Lys Val Phe Leu Thr Val Pro Ser Leu Ser Ser Thr Ala Glu
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 Glu Lys Phe Ile Lys Lys Gly Glu Phe Ala Gly Asp Asp Ser Leu Leu
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 Lys His Ile Tyr Asn Met Asp Pro Ser Lys Ser Val Pro Cys Ala Arg
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 Asp Lys Leu Ala Phe Thr Gln Ser Arg Ala Ala Ser Tyr Phe Phe Asp
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 Gly Gln Val Thr Arg Phe Asp Ile Glu Ile Arg Thr Pro Ala Asp Asn
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 Gly Pro Val His Leu Glu Asp Thr Leu Lys Lys Ala Gln Ile Asn Asp
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Pro Lys Thr Asn Lys Gln Gly Glu Lys Ser Lys Asp Ala Pro Ser Trp	1395	1400	1405
Asp Pro Ile Gly Leu Lys Phe Leu Glu Gln Lys Ala Pro Arg Asp Ser	1410	1415	1420
His Cys His Leu Ser Ser Ser Pro Arg Ala Ile Glu His Ala Tyr Gln	1425	1430	1435
Tyr Gly Gly Thr Ala Asn Ser Arg Gln Glu Phe Glu His Glu Gln Gly	1445	1450	1455
Asp Phe Gly Glu Lys Ser Gln Phe Ala Ile Arg Leu Lys Thr Arg Ser	1460	1465	1470
Ser His Gly Met Ile Phe Tyr Val Ser Asp Gln Glu Glu Asn Asp Phe	1475	1480	1485
Met Thr Leu Phe Leu Ala His Gly Arg Leu Val Phe Met Phe Asn Val	1490	1495	1500
Gly His Lys Lys Leu Lys Ile Arg Ser Gln Glu Lys Tyr Asn Asp Gly	1505	1510	1515
Leu Trp His Asp Val Ile Phe Ile Arg Glu Lys Ser Ser Gly Arg Leu	1525	1530	1535
Val Ile Asp Gly Leu Arg Val Leu Glu Glu Arg Leu Pro Pro Ser Gly	1540	1545	1550
Ala Ala Trp Lys Ile Lys Gly Pro Ile Tyr Leu Gly Gly Val Ala Pro	1555	1560	1565
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Gly Cys Leu Gly Asn Leu Gln Leu Asn Gly Ala Ser Ile Thr Ser Ala	1585	1590	1595
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Gly Thr Tyr Phe Ser Thr Glu Gly Gly Tyr Val Val Leu Asp Glu Ser	1620	1625	1630
Phe Asn Ile Gly Leu Lys Phe Glu Ile Ala Phe Glu Val Arg Pro Arg	1635	1640	1645
Ser Ser Ser Gly Thr Leu Val His Gly His Ser Val Asn Gly Glu Tyr	1650	1655	1660
Leu Asn Val His Met Arg Asn Gly Gln Val Ile Val Lys Val Asn Asn	1665	1670	1675
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Asp Gly Arg Trp His Arg Ile Thr Val Ile Arg Asp Ser Asn Val Val
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Gln Leu Asp Val Asp Ser Glu Val Asn His Val Val Gly Pro Leu Asn
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Glu Ser Leu Leu Thr Pro Arg Leu Ala Pro Ser Lys Pro Phe Thr Gly
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atg ggg ctt ctc cag ttg cta gct ttc agt ttc tta gcc ctg tgc aga 165
 Met Gly Leu Leu Gln Leu Leu Ala Phe Ser Phe Leu Ala Leu Cys Arg
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 Gly Ser Cys Tyr Pro Ala Thr Gly Asp Leu Leu Ile Gly Arg Ala Gln
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 Lys Leu Ser Val Thr Ser Thr Cys Gly Leu His Lys Pro Glu Pro Tyr
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 Cys Ile Val Ser His Leu Gln Glu Asp Lys Lys Cys Phe Ile Cys Asn
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Glu	Asn	Val	Val	Thr	Thr	Phe	Ala	Pro	Asn	Arg	Leu	Lys	Ile	Trp	Trp	
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Gln	Ser	Glu	Asn	Gly	Val	Glu	Asn	Val	Thr	Ile	Gln	Leu	Asp	Leu	Glu	
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Ala	Glu	Phe	His	Phe	Thr	His	Leu	Ile	Met	Thr	Phe	Lys	Thr	Phe	Arg	
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Pro	Ala	Ala	Met	Leu	Ile	Glu	Arg	Ser	Ser	Asp	Phe	Gly	Lys	Thr	Trp	
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Gly	Val	Tyr	Arg	Tyr	Phe	Ala	Tyr	Asp	Cys	Glu	Ala	Ser	Phe	Pro	Gly	
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Ile	Ser	Thr	Gly	Pro	Met	Lys	Lys	Val	Asp	Asp	Ile	Ile	Cys	Asp	Ser	
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Ala	Leu	Asp	Pro	Ala	Phe	Lys	Ile	Glu	Asp	Pro	Tyr	Ser	Pro	Arg	Ile	
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Gln	Asn	Leu	Leu	Lys	Ile	Thr	Asn	Leu	Arg	Ile	Lys	Phe	Val	Lys	Leu	
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Cys	Tyr	Gly	His	Ala	Ser	Glu	Cys	Ala	Pro	Val	Asp	Gly	Phe	Asn	Glu	
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Glu	Val	Glu	Gly	Met	Val	His	Gly	His	Cys	Met	Cys	Arg	His	Asn	Thr	
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Lys	Gly	Leu	Asn	Cys	Glu	Leu	Cys	Met	Asp	Phe	Tyr	His	Asp	Leu	Pro	
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Trp	Arg	Pro	Ala	Glu	Gly	Arg	Asn	Ser	Asn	Ala	Cys	Lys	Lys	Cys	Asn	
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Cys	Asn	Glu	His	Ser	Ile	Ser	Cys	His	Phe	Asp	Met	Ala	Val	Tyr	Leu	
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Phe	Ser	Thr	Gly	Leu	Ile	Ala	Gly	Gln	Cys	Arg	Cys	Lys	Leu	Asn	Val	
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His	Trp	Gly	Leu	Ser	Asn	Asp	Leu	Asp	Gly	Cys	Arg	Pro	Cys	Asp	Cys	
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Gln	Asp	Arg	Ile	Pro	Ser	Trp	Thr	Gly	Ala	Gly	Phe	Val	Arg	Val	Pro					
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Cys Asn Pro Val Thr Gly Gln Cys His Cys Phe Gln Gly Val Tyr Ala	
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Gln Cys His Asn Asn Ile Asp Thr Thr Asp Pro Glu Ala Cys Asp Lys	
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Arg Gly Ile Glu Thr Pro Gln Cys Asp Gln Ser Thr Gly Gln Cys Val	
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Cys Val Glu Gly Val Glu Gly Pro Arg Cys Asp Lys Cys Thr Arg Gly	
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Tyr Ser Gly Val Phe Pro Asp Cys Thr Pro Cys His Gln Cys Phe Ala	
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Leu Trp Asp Val Ile Ile Ala Glu Leu Thr Asn Arg Thr His Arg Phe	
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Arg Glu Thr Val Asp Ser Val Glu Arg Lys Val Ser Glu Ile Lys Asp	
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Ile Leu Ala Gln Ser Pro Ala Ala Glu Pro Leu Lys Asn Ile Gly Asn	
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ctc ttt gag gaa gca gag aaa ctg att aaa gat gtt aca gaa atg atg	3909
Leu Phe Glu Glu Ala Glu Lys Leu Ile Lys Asp Val Thr Glu Met Met	
1250 1255 1260	
gct caa gta gaa gtg aaa tta tct gac aca act tcc caa agc aac agc	3957
Ala Gln Val Glu Val Lys Leu Ser Asp Thr Thr Ser Gln Ser Asn Ser	
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aca gcc aaa gaa ctg gat tct cta cag aca gaa gcc gaa agc cta gac	4005
Thr Ala Lys Glu Leu Asp Ser Leu Gln Thr Glu Ala Glu Ser Leu Asp	
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Asn Thr Val Lys Glu Leu Ala Glu Gln Leu Glu Phe Ile Lys Asn Ser	
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Leu Glu Ala Glu Glu Arg Val Asn Ala Ser Thr Thr Glu Pro Asn Ser	
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act gtg gag cag tca gcc ctc atg aga gac aga gta gaa gac gtg atg	4197
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Ala Ala Glu Met Thr Cys Gly Thr Pro Pro Gly Ala Ser Cys Ser Glu	
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Cys Gly Gly Pro Gly Cys Gly Gly Leu Val Thr Val Ala His Asn Ala	
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Trp Gln Lys Ala Met Asp Leu Asp Gln Asp Val Leu Ser Ala Leu Ala	
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Glu Val Glu Gln Leu Ser Lys Met Val Ser Glu Ala Lys Leu Arg Ala	
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Asp Glu Ala Lys Gln Ser Ala Glu Asp Ile Leu Leu Lys Thr Asn Ala	
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Pro Gln Gln Leu Gln Asn Leu Thr Glu Asp Ile Arg Glu Arg Val Glu	

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gat gaa gac att caa gga acc cag aac ctg tta act tcg att gag tct Asp Glu Asp Ile Gln Gly Thr Gln Asn Leu Leu Thr Ser Ile Glu Ser 1620	1625	1630	5013
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aac tcc ggg gag gca gaa tat att gaa aaa gta gta tat act gtg aag Asn Ser Gly Glu Ala Glu Tyr Ile Glu Lys Val Val Tyr Thr Val Lys 1665	1670	1675	5157
caa agt gca gaa gat gtt aag aag act tta gat ggt gaa ctt gat gaa Gln Ser Ala Glu Asp Val Lys Lys Thr Leu Asp Gly Glu Leu Asp Glu 1685	1690	1695	5205
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tta gca aga ctg gaa gga gaa gtc cgt tca ctc cta aag gat ata agc Leu Ala Arg Leu Glu Gly Glu Val Arg Ser Leu Leu Lys Asp Ile Ser 1765	1770	1775	5445
cag aaa gtt gct gtg tat agc aca tgc ttg taacagagga gaataaaaaa Gln Lys Val Ala Val Tyr Ser Thr Cys Leu 1780	1785		5495

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 Ser Gln Asp Pro Tyr His Glu Thr Leu Asn Pro Asp Ser His Leu Ile
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 Glu Asn Val Val Thr Thr Phe Ala Pro Asn Arg Leu Lys Ile Trp Trp
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 Arg Tyr Ser Asp Ile Glu Pro Ser Thr Glu Gly Glu Val Ile Phe Arg
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 Ala Leu Asp Pro Ala Phe Lys Ile Glu Asp Pro Tyr Ser Pro Arg Ile
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 Gln Asn Leu Leu Lys Ile Thr Asn Leu Arg Ile Lys Phe Val Lys Leu
 225 230 235 240
 His Thr Leu Gly Asp Asn Leu Leu Asp Ser Arg Met Glu Ile Arg Glu
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 Lys Tyr Tyr Tyr Ala Val Tyr Asp Met Val Val Arg Gly Asn Cys Phe

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Glu	Val	Glu	Gly	Met	Val	His	Gly	His	Cys	Met	Cys	Arg	His	Asr	Thr		
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Cys	Asn	Glu	His	Ser	Ile	Ser	Cys	His	Phe	Asp	Met	Ala	Val	Tyr	Leu		
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 Glu Lys Ala Val Ile Thr Val Gln Arg Pro Gly Arg Ile Pro Thr Ser
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Cys Gly Gly Pro Gly Cys Gly Gly Leu Val Thr Val Ala His Asn Ala 1425	1430	1435 1440
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Glu Val Glu Gln Leu Ser Lys Met Val Ser Glu Ala Lys Leu Arg Ala 1460	1465	1470
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Lys Gln Ile Arg Asn Phe Leu Thr Gln Asp Ser Ala Asp Leu Asp Ser 1505	1510	1515 1520
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Cys Glu Gln Cys Lys Pro Phe Tyr Tyr Gln His Pro Glu Arg Asp Ile	
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Arg Asp Pro Asn Phe Cys Glu Arg Cys Thr Cys Asp Pro Ala Gly Ser	
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Gln Asn Glu Gly Ile Cys Asp Ser Tyr Thr Asp Phe Ser Thr Gly Leu	
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Ile Ala Gly Gln Cys Arg Cys Lys Leu Asn Val Glu Gly Glu His Cys	
405 410 415	
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Asp Val Cys Lys Glu Gly Phe Tyr Asp Leu Ser Ser Glu Asp Pro Phe	
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Gly Cys Lys Ser Cys Ala Cys Asn Pro Leu Gly Thr Ile Pro Gly Gly	
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Asn Pro Cys Asp Ser Glu Thr Gly His Cys Tyr Cys Lys Arg Leu Val	
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aca gga cag cat tgt gac cag tgc ctg cca gag cac tgg ggc tta agc	1440
Thr Gly Gln His Cys Asp Gln Cys Leu Pro Glu His Trp Gly Leu Ser	
465 470 475 480	
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Asn Asp Leu Asp Gly Cys Arg Pro Cys Asp Cys Asp Leu Gly Gly Ala	
485 490 495	
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Leu Asn Asn Ser Cys Phe Ala Glu Ser Gly Gln Cys Ser Cys Arg Pro	
500 505 510	
cac atg att gga cgt cag tgc aac gaa gtg gaa cct ggt tac tac ttt	1584

His	Met	Ile	Gly	Arg	Gln	Cys	Asn	Glu	Val	Glu	Pro	Gly	Tyr	Tyr	Phe	
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gcc	acc	ctg	gat	cac	tac	ctc	tat	gaa	gcg	gag	gaa	gcc	aac	ttg	ggg	1632
Ala	Thr	Leu	Asp	His	Tyr	Leu	Tyr	Glu	Ala	Glu	Glu	Ala	Asn	Leu	Gly	
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cct	ggg	gtt	agc	ata	gtg	gag	cgg	caa	tat	atc	cag	gac	cgg	att	ccc	1680
Pro	Gly	Val	Ser	Ile	Val	Glu	Arg	Gln	Tyr	Ile	Gln	Asp	Arg	Ile	Pro	
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Ser	Trp	Thr	Gly	Ala	Gly	Phe	Val	Arg	Val	Pro	Glu	Gly	Ala	Tyr	Leu	
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Glu	Phe	Phe	Ile	Asp	Asn	Ile	Pro	Tyr	Ser	Met	Glu	Tyr	Asp	Ile	Leu	
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Ile	Arg	Tyr	Glu	Pro	Gln	Leu	Pro	Asp	His	Trp	Glu	Lys	Ala	Val	Ile	
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Thr	Val	Gln	Arg	Pro	Gly	Arg	Ile	Pro	Thr	Ser	Ser	Arg	Cys	Gly	Asn	
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Thr	Ile	Pro	Asp	Asp	Asp	Asn	Gln	Val	Val	Ser	Leu	Ser	Pro	Gly	Ser	
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Arg	Tyr	Val	Val	Leu	Pro	Arg	Pro	Val	Cys	Phe	Glu	Lys	Gly	Thr	Asn	
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tac	acg	gtg	agg	ttg	gag	ctg	cct	cag	tac	acc	tcc	tct	gat	agc	gac	2016
Tyr	Thr	Val	Arg	Leu	Glu	Leu	Pro	Gln	Tyr	Thr	Ser	Ser	Asp	Ser	Asp	
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gtg	gag	agc	ccc	tac	acg	ctg	atc	gat	tct	ctt	gtt	ctc	atg	cca	tac	2064
Val	Glu	Ser	Pro	Tyr	Thr	Leu	Ile	Asp	Ser	Leu	Val	Leu	Met	Pro	Tyr	
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tgt	aaa	tca	ctg	gac	atc	ttc	acc	gtg	gga	ggg	tca	gga	gat	ggg	gtg	2112
Cys	Lys	Ser	Leu	Asp	Ile	Phe	Thr	Val	Gly	Gly	Ser	Gly	Asp	Gly	Val	
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gtc	acc	aac	agt	gcc	tgg	gaa	acc	ttt	cag	aga	tac	cga	tgt	cta	gag	2160
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Asn	Ser	Arg	Ser	Val	Val	Lys	Thr	Pro	Met	Thr	Asp	Val	Cys	Arg	Asn	
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atc	atc	ttt	agc	att	tct	gcc	ctg	tta	cac	cag	aca	ggc	ctg	gct	tgt	2256
Ile	Ile	Phe	Ser	Ile	Ser	Ala	Leu	Leu	His	Gln	Thr	Gly	Leu	Ala	Cys	
			740					745					750			
gaa	tgc	gac	cct	cag	ggg	tcg	tta	agt	tcc	gtg	tgt	gat	ccc	aac	gga	2304
Glu	Cys	Asp	Pro	Gln	Gly	Ser	Leu	Ser	Ser	Val	Cys	Asp	Pro	Asn	Gly	

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tgt gca cct gga act ttt ggc ttt ggc ccc agt gga tgc aaa cct tgt Cys Ala Pro Gly Thr Phe Gly Phe Gly Pro Ser Gly Cys Lys Pro Cys 785 790 795 800			2400
gag tgc cat ctg caa gga tct gtc aat gcc ttc tgc aat ccc gtc act Glu Cys His Leu Gln Gly Ser Val Asn Ala Phe Cys Asn Pro Val Thr 805 810 815			2448
ggc cag tgc cac tgt ttc cag gga gtg tat gct cgg cag tgt gat cgg Gly Gln Cys His Cys Phe Gln Gly Val Tyr Ala Arg Gln Cys Asp Arg 820 825 830			2496
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aat ggc cac gcc gat gac tgc gac cca gtg act ggg gag tgc ttg aac Asn Gly His Ala Asp Asp Cys Asp Pro Val Thr Gly Glu Cys Leu Asn 850 855 860			2592
tgc cag gac tac acc atg ggt cat aac tgt gaa agg tgc ttg gct ggt Cys Gln Asp Tyr Thr Met Gly His Asn Cys Glu Arg Cys Leu Ala Gly 865 870 875 880			2640
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tac att ggt tcc aga tgt gac gac tgt gcc tca gga tac ttt ggc aat Tyr Ile Gly Ser Arg Cys Asp Asp Cys Ala Ser Gly Tyr Phe Gly Asn 930 935 940			2832
cca tca gaa gtt ggg ggg tcg tgt cag cct tgc cag tgt cac aac aac Pro Ser Glu Val Gly Gly Ser Cys Gln Pro Cys Gln Cys His Asn Asn 945 950 955 960			2880
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ctc aag tgc ctg tac cac acg gaa ggg gaa cac tgt cag ttc tgc cgg Leu Lys Cys Leu Tyr His Thr Glu Gly Glu His Cys Gln Phe Cys Arg 980 985 990			2976
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Cys Asn Tyr Leu Gly Thr Val Gln Glu His Cys Asn Gly Ser Asp Cys	
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1125 1130 1135	
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Glu Gly Pro Arg Cys Asp Lys Cys Thr Arg Gly Tyr Ser Gly Val Phe	
1140 1145 1150	
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Pro Asp Cys Thr Pro Cys His Gln Cys Phe Ala Leu Trp Asp Val Ile	
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Ile Ala Glu Leu Thr Asn Arg Thr His Arg Phe Leu Glu Lys Ala Lys	
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Ala Leu Lys Ile Ser Gly Val Ile Gly Pro Tyr Arg Glu Thr Val Asp	
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Ser Val Glu Arg Lys Val Ser Glu Ile Lys Asp Ile Leu Ala Gln Ser	
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Pro Ala Ala Glu Pro Leu Lys Asn Ile Gly Asn Leu Phe Glu Glu Ala	
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Glu Lys Leu Ile Lys Asp Val Thr Glu Met Met Ala Gln Val Glu Val	
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Lys	Ala	Glu	Met	Leu	Gln	Asn	Glu	Ala	Lys	Thr	Leu	Leu	Ala	Gln	Ala		
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1730

1735

1740

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Leu Gln Glu Asp Lys Lys Cys Phe Ile Cys Asn Ser Gln Asp Pro Tyr
 50 55 60

His Glu Thr Leu Asn Pro Asp Ser His Leu Ile Glu Asn Val Val Thr
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Thr Phe Ala Pro Asn Arg Leu Lys Ile Trp Trp Gln Ser Glu Asn Gly
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Val Glu Asn Val Thr Ile Gln Leu Asp Leu Glu Ala Glu Phe His Phe
 100 105 110

Thr His Leu Ile Met Thr Phe Lys Thr Phe Arg Pro Ala Ala Met Leu
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Ile Glu Arg Ser Ser Asp Phe Gly Lys Thr Trp Gly Val Tyr Arg Tyr
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Met Lys Lys Val Asp Asp Ile Ile Cys Asp Ser Arg Tyr Ser Asp Ile
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Glu Pro Ser Thr Glu Gly Glu Val Ile Phe Arg Ala Leu Asp Pro Ala
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Phe Lys Ile Glu Asp Pro Tyr Ser Pro Arg Ile Gln Asn Leu Leu Lys
 195 200 205

Ile Thr Asn Leu Arg Ile Lys Phe Val Lys Leu His Thr Leu Gly Asp

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Cys Glu Gln Cys Lys Pro Phe Tyr Tyr Gln His Pro Glu Arg Asp Ile 355 360 365		
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Ile Ala Gly Gln Cys Arg Cys Lys Leu Asn Val Glu Gly Glu His Cys 405 410 415		
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Asn Pro Cys Asp Ser Glu Thr Gly His Cys Tyr Cys Lys Arg Leu Val 450 455 460		
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His Met Ile Gly Arg Gln Cys Asn Glu Val Glu Pro Gly Tyr Tyr Phe 515 520 525		
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 675 680 685
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Thr Val Glu Gln Ser Ala Leu Thr Arg Asp Arg Val Glu Asp Leu Met	
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ttg gag cga gag tct ccg ttc aag gag cag cag gag gaa cag gca cgc	4305
Leu Glu Arg Glu Ser Pro Phe Lys Glu Gln Gln Glu Glu Gln Ala Arg	
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Leu Leu Asp Glu Leu Ala Gly Lys Leu Gln Ser Leu Asp Leu Ser Ala	
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gct gca cag atg acc tgt gga aca cct cca ggg gct gac tgt tct gaa	4401
Ala Ala Gln Met Thr Cys Gly Thr Pro Pro Gly Ala Asp Cys Ser Glu	
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agt gaa tgt ggt ggc ccc aac tgc aga act gac gaa gga gag aag aag	4449
Ser Glu Cys Gly Gly Pro Asn Cys Arg Thr Asp Glu Gly Glu Lys Lys	
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Cys Gly Gly Pro Gly Cys Gly Gly Leu Val Thr Val Ala His Ser Ala	
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Trp Gln Lys Ala Met Asp Phe Asp Arg Asp Val Leu Ser Ala Leu Ala	
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gaa gtc gaa cag ctc tcc aag atg gtc tct gaa gca aaa gtg aga gca	4593
Glu Val Glu Gln Leu Ser Lys Met Val Ser Glu Ala Lys Val Arg Ala	
	1460 1465 1470
gat gag gcg aag cag aat gcg cag gat gtc ctg tta aaa aca aat gct	4641
Asp Glu Ala Lys Gln Asn Ala Gln Asp Val Leu Leu Lys Thr Asn Ala	
	1475 1480 1485
acc aaa gaa aaa gtg gac aag agc aac gag gac ctg cgg aac ctc atc	4689
Thr Lys Glu Lys Val Asp Lys Ser Asn Glu Asp Leu Arg Asn Leu Ile	
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Lys Gln Ile Arg Asn Phe Leu Thr Glu Asp Ser Ala Asp Leu Asp Ser	
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Ile Glu Ala Val Ala Asn Glu Val Leu Lys Ser Gly Asn Ala Ser Thr	
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cca cag cag tta cag aac cta aca gaa gac att cgg gag cga gtt gaa	4833
Pro Gln Gln Leu Gln Asn Leu Thr Glu Asp Ile Arg Glu Arg Val Glu	
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gcc aga gct gag ctg ttg ctt gag gaa gct aag aga gca agc aaa agt	4929
Ala Arg Ala Glu Leu Leu Leu Glu Glu Ala Lys Arg Ala Ser Lys Ser	
	1570 1575 1580
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Ala Thr Asp Val Lys Val Thr Ala Asp Met Val Lys Glu Ala Leu Glu	
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Glu Thr Ala Ala Ser Glu Glu Thr Leu Thr Asn Ala Ser Gln Arg Ile	
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Ser Lys Leu Glu Arg Asn Val Glu Glu Leu Lys Arg Lys Ala Ala Gln	
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Gln Asn Ala Asp Asp Val Lys Lys Thr Leu Asp Gly Glu Leu Asp Glu				
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Lys Tyr Lys Lys Val Glu Ser Leu Ile Ala Gln Lys Thr Glu Glu Ser				
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gca gat gcc agg agg aaa gct gag ctg cta caa aat gaa gca aaa aca				5361
Ala Asp Ala Arg Arg Lys Ala Glu Leu Leu Gln Asn Glu Ala Lys Thr				
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ctc ttg gct caa gct aac agc aag ctc cag ctg ttg gaa gac tta gaa				5409
Leu Leu Ala Gln Ala Asn Ser Lys Leu Gln Leu Leu Glu Asp Leu Glu				
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Arg Lys Tyr Glu Asp Asn Gln Lys Tyr Leu Glu Asp Lys Ala Gln Glu				
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Glu Lys Val Ala Val Tyr Ser Thr Cys Leu				
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Cys Ile Val Ser His Leu Gln Glu Asp Lys Lys Cys Phe Ile Cys Asp	
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 Gln Ser Glu Asn Gly Val Glu Asn Val Thr Ile Gln Leu Asp Leu Glu
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 Gly Val Tyr Arg Tyr Phe Ala Tyr Asp Cys Glu Ser Ser Phe Pro Gly
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 Ile Ser Thr Gly Pro Met Lys Lys Val Asp Asp Ile Ile Cys Asp Ser
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 Arg Tyr Ser Asp Ile Glu Pro Ser Thr Glu Gly Glu Val Ile Phe Arg
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 Ala Leu Asp Pro Ala Phe Lys Ile Glu Asp Pro Tyr Ser Pro Arg Ile
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 Gln Asn Leu Leu Lys Ile Thr Asn Leu Arg Ile Lys Phe Val Lys Leu
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 His Thr Leu Gly Asp Asn Leu Leu Asp Ser Arg Met Glu Ile Arg Glu
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 Lys Tyr Tyr Tyr Ala Val Tyr Asp Met Val Val Arg Gly Asn Cys Phe
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 Cys Tyr Gly His Ala Ser Glu Cys Ala Pro Val Asp Gly Val Asn Glu
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 Glu Val Glu Gly Met Val His Gly His Cys Met Cys Arg His Asn Thr
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 Lys Gly Leu Asn Cys Glu Leu Cys Met Asp Phe Tyr His Asp Leu Pro
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 Trp Arg Pro Ala Glu Gly Arg Asn Ser Asn Ala Cys Lys Lys Cys Asn
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 Cys Asn Glu His Ser Ser Ser Cys His Phe Asp Met Ala Val Phe Leu
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 Arg Cys Leu Ala Gly Tyr Tyr Gly Asp Pro Ile Ile Gly Ser Gly Asp
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 Ile Cys Asp Ser Arg Asp Pro Tyr His Glu Thr Leu Asn Pro Asp Ser
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 His Leu Ile Glu Asn Val Val Thr Thr Phe Ala Pro Asn Arg Leu Lys
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 Ile Trp Trp Gln Ser Glu Asn Gly Val Glu Asn Val Thr Ile Gln Leu
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Tyr Cys Tyr Cys Lys Arg Leu Val Thr Gly Gln Arg Cys Asp Gln Cys							
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Ser Gly Gln Cys Ser Cys Leu Pro His Met Ile Gly Arg Gln Cys Asn							
	465	470	475	480			
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Glu Val Glu Ser Gly Tyr Tyr Phe Thr Thr Leu Asp His Tyr Ile Tyr							
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gaa gcc gag gaa gcc aat ctg ggg cct gga gtc gtt gtg gtg gaa agg	1536						
Glu Ala Glu Glu Ala Asn Leu Gly Pro Gly Val Val Val Val Glu Arg							
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cag tac att cag gac cgc att cct tcc tgg aca gga cct ggc ttc gtc	1584						
Gln Tyr Ile Gln Asp Arg Ile Pro Ser Trp Thr Gly Pro Gly Phe Val							
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cgg gtg cct gaa ggg gct tat ttg gag ttt ttc att gac aac ata cca	1632						
Arg Val Pro Glu Gly Ala Tyr Leu Glu Phe Phe Ile Asp Asn Ile Pro							
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tat tcc atg gag tat gaa atc ctg att cgc tat gag cca cag ctg ccg	1680						
Tyr Ser Met Glu Tyr Glu Ile Leu Ile Arg Tyr Glu Pro Gln Leu Pro							
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gac cac tgg gag aaa gct gtc atc act gta cag cgg ccg ggg aag att	1728						
Asp His Trp Glu Lys Ala Val Ile Thr Val Gln Arg Pro Gly Lys Ile							
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cca gcc agc agc cga tgt ggt aac acc gtt ccc gat gat gac aac cag	1776						
Pro Ala Ser Ser Arg Cys Gly Asn Thr Val Pro Asp Asp Asp Asn Gln							
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Val Val Ser Leu Ser Pro Gly Ser Arg Tyr Val Val Leu Pro Arg Pro							
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Val Cys Phe Glu Lys Gly Met Asn Tyr Thr Val Arg Leu Glu Leu Pro							
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Gln Tyr Thr Ala Ser Gly Ser Asp Val Glu Ser Pro Tyr Thr Phe Ile							
	625	630	635	640			

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Asp Ser Leu Val Leu Met Pro Tyr Cys Lys Ser Leu Asp Ile Phe Thr	
645 650 655	
gtt ggc ggc tca ggc gat ggg gag gtc acc aat agt gcc tgg gaa acc	2016
Val Gly Gly Ser Gly Asp Gly Glu Val Thr Asn Ser Ala Trp Glu Thr	
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Phe Gln Arg Tyr Arg Cys Leu Glu Asn Ser Arg Ser Val Val Lys Thr	
675 680 685	
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Pro Met Thr Asp Val Cys Arg Asn Ile Ile Phe Ser Ile Ser Ala Leu	
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Ile His Gln Thr Gly Leu Ala Cys Glu Cys Asp Pro Gln Gly Ser Leu	
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agt tct gtg tgt gac ccc aat ggt ggc cag tgc cag tgc cgt cct aat	2208
Ser Ser Val Cys Asp Pro Asn Gly Gly Gln Cys Gln Cys Arg Pro Asn	
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Val Val Gly Arg Thr Cys Asn Arg Cys Ala Pro Gly Thr Phe Gly Phe	
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Gly Pro Asn Gly Cys Lys Pro Cys Asp Cys His Leu Gln Gly Ser Ala	
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Ser Ala Phe Cys Asp Ala Ile Thr Gly Gln Cys His Cys Phe Gln Gly	
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Ile Tyr Ala Arg Gln Cys Asp Arg Cys Leu Pro Gly Tyr Trp Gly Phe	
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Pro Ser Cys Gln Pro Cys Gln Cys Asn Gly His Ala Leu Asp Cys Asp	
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Asn Cys Glu Arg Cys Leu Ala Gly Tyr Tyr Gly Asp Pro Ile Ile Gly	
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Ser Gly Asp His Cys Arg Pro Cys Pro Cys Pro Asp Gly Pro Asp Ser	
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Gly Arg Gln Phe Ala Arg Ser Cys Tyr Gln Asp Pro Val Thr Leu Gln	
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Leu Ala Cys Val Cys Asp Pro Gly Tyr Ile Gly Ser Arg Cys Asp Asp	
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Cys Ala Ser Gly Phe Phe Gly Asn Pro Ser Asp Phe Gly Gly Ser Cys	
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Gln Pro Cys Gln Cys His His Asn Ile Asp Thr Thr Asp Pro Glu Ala	
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Cys Asp Lys Asp Thr Gly Arg Cys Leu Lys Cys Leu Tyr His Thr Glu	
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Gly Asp His Cys Gln Leu Cys Gln Tyr Gly Tyr Tyr Gly Asp Ala Leu	
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Cys Ser Cys Leu Pro Asn Val Ile Gly Gln Asn Cys Asp Arg Cys Ala	
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Gln Glu Leu Phe Trp Gly Asp Pro Asp Val Glu Cys Arg Ala Cys Asp	
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Cys Asp Pro Arg Gly Ile Glu Thr Pro Gln Cys Asp Gln Ser Thr Gly	
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Gln Cys Val Cys Val Glu Gly Val Glu Gly Pro Arg Cys Asp Lys Cys	
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Thr Arg Gly Tyr Ser Gly Val Phe Pro Asp Cys Thr Pro Cys His Gln	
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Cys	Phe	Ala	Leu	Trp	Asp	Ala	Ile	Ile	Gly	Glu	Leu	Thr	Asn	Arg	Thr	
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cac	aaa	ttc	ctg	gag	aaa	gcc	aag	gct	ctg	aaa	atc	agt	ggg	gtg	att	3456
His	Lys	Phe	Leu	Glu	Lys	Ala	Lys	Ala	Leu	Lys	Ile	Ser	Gly	Val	Ile	
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Gly	Pro	Tyr	Arg	Glu	Thr	Val	Asp	Ser	Val	Glu	Lys	Lys	Val	Asn	Glu	
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Ile	Lys	Asp	Ile	Leu	Ala	Gln	Ser	Pro	Ala	Ala	Glu	Pro	Leu	Lys	Asn	
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Ile	Gly	Ile	Leu	Phe	Glu	Glu	Ala	Glu	Lys	Leu	Thr	Lys	Asp	Val	Thr	
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Glu	Lys	Met	Ala	Gln	Val	Glu	Val	Lys	Leu	Thr	Asp	Thr	Ala	Ser	Gln	
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Ser	Asn	Ser	Thr	Ala	Gly	Glu	Leu	Gly	Ala	Leu	Gln	Ala	Glu	Ala	Glu	
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Ser	Leu	Asp	Lys	Thr	Val	Lys	Glu	Leu	Ala	Glu	Gln	Leu	Glu	Phe	Ile	
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Lys	Asn	Ser	Asp	Ile	Gln	Gly	Ala	Leu	Asp	Ser	Ile	Thr	Lys	Tyr	Phe	
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Gln	Met	Ser	Leu	Glu	Ala	Glu	Lys	Arg	Val	Asn	Ala	Ser	Thr	Thr	Asp	
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Pro	Asn	Ser	Thr	Val	Glu	Gln	Ser	Ala	Leu	Thr	Arg	Asp	Arg	Val	Glu	
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Asp	Leu	Met	Leu	Glu	Arg	Glu	Ser	Pro	Phe	Lys	Glu	Gln	Gln	Glu	Glu	
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Le																

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His Ser Ala Trp Gln Lys Ala Met Asp Phe Asp Arg Asp Val Leu Ser			
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gcc ctg gct gaa gtc gaa cag ctc tcc aag atg gtc tct gaa gca aaa			4224
Ala Leu Ala Glu Val Glu Gln Leu Ser Lys Met Val Ser Glu Ala Lys			
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Val Arg Ala Asp Glu Ala Lys Gln Asn Ala Gln Asp Val Leu Leu Lys			
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Thr Asn Ala Thr Lys Glu Lys Val Asp Lys Ser Asn Glu Asp Leu Arg			
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Gln Arg Ile Ser Lys Leu Glu Arg Asn Val Glu Glu Leu Lys Arg Lys			
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gct gcc cag aac tct ggg gag gca gaa tat atc gaa aaa gta gta tat			4848
Ala Ala Gln Asn Ser Gly Glu Ala Glu Tyr Ile Glu Lys Val Val Tyr			
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 35 40 45
 Ile Trp Trp Gln Ser Glu Asn Gly Val Glu Asn Val Thr Ile Gln Leu
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 Asp Leu Glu Ala Glu Phe His Phe Thr His Leu Ile Met Thr Phe Lys
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 Thr Phe Arg Pro Ala Ala Met Leu Ile Glu Arg Ser Ser Asp Phe Gly
 85 90 95
 Lys Thr Trp Gly Val Tyr Arg Tyr Phe Ala Tyr Asp Cys Glu Ser Ser

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1075	1080	1085
Gln Cys Val Cys Val Glu Gly Val Glu Gly Pro Arg Cys Asp Lys Cys 1090	1095	1100
Thr Arg Gly Tyr Ser Gly Val Phe Pro Asp Cys Thr Pro Cys His Gln 1105	1110	1115 1120
Cys Phe Ala Leu Trp Asp Ala Ile Ile Gly Glu Leu Thr Asn Arg Thr 1125	1130	1135
His Lys Phe Leu Glu Lys Ala Lys Ala Leu Lys Ile Ser Gly Val Ile 1140	1145	1150
Gly Pro Tyr Arg Glu Thr Val Asp Ser Val Glu Lys Lys Val Asn Glu 1155	1160	1165
Ile Lys Asp Ile Leu Ala Gln Ser Pro Ala Ala Glu Pro Leu Lys Asn 1170	1175	1180
Ile Gly Ile Leu Phe Glu Glu Ala Glu Lys Leu Thr Lys Asp Val Thr 1185	1190	1195 1200
Glu Lys Met Ala Gln Val Glu Val Lys Leu Thr Asp Thr Ala Ser Gln 1205	1210	1215
Ser Asn Ser Thr Ala Gly Glu Leu Gly Ala Leu Gln Ala Glu Ala Glu 1220	1225	1230
Ser Leu Asp Lys Thr Val Lys Glu Leu Ala Glu Gln Leu Glu Phe Ile 1235	1240	1245
Lys Asn Ser Asp Ile Gln Gly Ala Leu Asp Ser Ile Thr Lys Tyr Phe 1250	1255	1260
Gln Met Ser Leu Glu Ala Glu Lys Arg Val Asn Ala Ser Thr Thr Asp 1265	1270	1275 1280
Pro Asn Ser Thr Val Glu Gln Ser Ala Leu Thr Arg Asp Arg Val Glu 1285	1290	1295
Asp Leu Met Leu Glu Arg Glu Ser Pro Phe Lys Glu Gln Gln Glu Glu 1300	1305	1310
Gln Ala Arg Leu Leu Asp Glu Leu Ala Gly Lys Leu Gln Ser Leu Asp 1315	1320	1325
Leu Ser Ala Ala Ala Gln Met Thr Cys Gly Thr Pro Pro Gly Ala Asp 1330	1335	1340
Cys Ser Glu Ser Glu Cys Gly Gly Pro Asn Cys Arg Thr Asp Glu Gly 1345	1350	1355 1360
Glu Lys Lys Cys Gly Gly Pro Gly Cys Gly Gly Leu Val Thr Val Ala 1365	1370	1375
His Ser Ala Trp Gln Lys Ala Met Asp Phe Asp Arg Asp Val Leu Ser 1380	1385	1390
Ala Leu Ala Glu Val Glu Gln Leu Ser Lys Met Val Ser Glu Ala Lys 1395	1400	1405

Val Arg Ala Asp Glu Ala Lys Gln Asn Ala Gln Asp Val Leu Leu Lys
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Thr Asn Ala Thr Lys Glu Lys Val Asp Lys Ser Asn Glu Asp Leu Arg
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Asn Leu Ile Lys Gln Ile Arg Asn Phe Leu Thr Glu Asp Ser Ala Asp
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Leu Asp Ser Ile Glu Ala Val Ala Asn Glu Val Leu Lys Ser Gly Asn
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Ala Ser Thr Pro Gln Gln Leu Gln Asn Leu Thr Glu Asp Ile Arg Glu
1475 1480 1485

Arg Val Glu Thr Leu Ser Gln Val Glu Val Ile Leu Gln Gln Ser Ala
1490 1495 1500

Ala Asp Ile Ala Arg Ala Glu Leu Leu Leu Glu Glu Ala Lys Arg Ala
1505 1510 1515 1520

Ser Lys Ser Ala Thr Asp Val Lys Val Thr Ala Asp Met Val Lys Glu
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Ala Leu Glu Glu Ala Glu Lys Ala Gln Val Ala Ala Glu Lys Ala Ile
1540 1545 1550

Lys Gln Ala Asp Glu Asp Ile Gln Gly Thr Gln Asn Leu Leu Thr Ser
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Ile Glu Ser Glu Thr Ala Ala Ser Glu Glu Thr Leu Thr Asn Ala Ser
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Gln Arg Ile Ser Lys Leu Glu Arg Asn Val Glu Glu Leu Lys Arg Lys
1585 1590 1595 1600

Ala Ala Gln Asn Ser Gly Glu Ala Glu Tyr Ile Glu Lys Val Val Tyr
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Ser Val Lys Gln Asn Ala Asp Asp Val Lys Lys Thr Leu Asp Gly Glu
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Leu Asp Glu Lys Tyr Lys Lys Val Glu Ser Leu Ile Ala Gln Lys Thr
1635 1640 1645

Glu Glu Ser Ala Asp Ala Arg Arg Lys Ala Glu Leu Leu Gln Asn Glu
1650 1655 1660

Ala Lys Thr Leu Leu Ala Gln Ala Asn Ser Lys Leu Gln Leu Leu Glu
1665 1670 1675 1680

Asp Leu Glu Arg Lys Tyr Glu Asp Asn Gln Lys Tyr Leu Glu Asp Lys
1685 1690 1695

Ala Gln Glu Leu Val Arg Leu Glu Gly Glu Val Arg Ser Leu Leu Lys
1700 1705 1710

Asp Ile Ser Glu Lys Val Ala Val Tyr Ser Thr Cys Leu
1715 1720 1725

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cgc ccg gag agc ttt gcc att tac aag cgc aca cgg gaa gac ggg ccc				772
Arg Pro Glu Ser Phe Ala Ile Tyr Lys Arg Thr Arg Glu Asp Gly Pro				
	160	165	170	
tgg att cct tac cag tac tac agt ggt tcc tgc gag aac acc tac tcc				820
Trp Ile Pro Tyr Gln Tyr Tyr Ser Gly Ser Cys Glu Asn Thr Tyr Ser				
	175	180	185	
aag gca aac cgc ggc ttc atc agg aca gga ggg gac gag cag cag gcc				868
Lys Ala Asn Arg Gly Phe Ile Arg Thr Gly Gly Asp Glu Gln Gln Ala				
	190	195	200	
ttg tgt act gat gaa ttc agt gac att tct ccc ctc act ggg ggc aac				916
Leu Cys Thr Asp Glu Phe Ser Asp Ile Ser Pro Leu Thr Gly Gly Asn				
	205	210	215	
gtg gcc ttt tct acc ctg gaa gga agg ccc agc gcc tat aac ttt gac				964
Val Ala Phe Ser Thr Leu Glu Gly Arg Pro Ser Ala Tyr Asn Phe Asp				
	220	225	230	235
aat agc cct gtg ctg cag gaa tgg gta act gcc act gac atc aga gta				1012
Asn Ser Pro Val Leu Gln Glu Trp Val Thr Ala Thr Asp Ile Arg Val				
	240	245	250	
act ctt aat cgc ctg aac act ttt gga gat gaa gtg ttt aac gat ccc				1060
Thr Leu Asn Arg Leu Asn Thr Phe Gly Asp Glu Val Phe Asn Asp Pro				
	255	260	265	
aaa gtt ctc aag tcc tat tat tat gcc atc tct gat ttt gct gta ggt				1108
Lys Val Leu Lys Ser Tyr Tyr Tyr Ala Ile Ser Asp Phe Ala Val Gly				
	270	275	280	
ggc aga tgt aaa tgt aat gga cac gca agc gag tgt atg aag aac gaa				1156
Gly Arg Cys Lys Cys Asn Gly His Ala Ser Glu Cys Met Lys Asn Glu				
	285	290	295	
ttt gat aag ctg gtg tgt aat tgc aaa cat aac aca tat gga gta gac				1204
Phe Asp Lys Leu Val Cys Asn Cys Lys His Asn Thr Tyr Gly Val Asp				
	300	305	310	315
tgt gaa aag tgt ctt cct ttc ttc aat gac cgg ccg tgg agg agg gca				1252
Cys Glu Lys Cys Leu Pro Phe Phe Asn Asp Arg Pro Trp Arg Arg Ala				
	320	325	330	
act gcg gaa agt gcc agt gaa tgc ctg ccc tgt gat tgc aat ggt cga				1300
Thr Ala Glu Ser Ala Ser Glu Cys Leu Pro Cys Asp Cys Asn Gly Arg				
	335	340	345	
tcc cag gaa tgc tac ttc gac cct gaa ctc tat cgt tcc act ggc cat				1348
Ser Gln Glu Cys Tyr Phe Asp Pro Glu Leu Tyr Arg Ser Thr Gly His				
	350	355	360	
ggg ggc cac tgt acc aac tgc cag gat aac aca gat ggc gcc cac tgt				1396
Gly Gly His Cys Thr Asn Cys Gln Asp Asn Thr Asp Gly Ala His Cys				
	365	370	375	
gag agg tgc cga gag aac ttc ttc cgc ctt ggc aac aat gaa gcc tgc				1444
Glu Arg Cys Arg Glu Asn Phe Phe Arg Leu Gly Asn Asn Glu Ala Cys				
	380	385	390	395

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Ser Ser Cys His Cys Ser Pro Val Gly Ser Leu Ser Thr Gln Cys Asp	
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agt tac ggc aga tgc agc tgt aag cca gga gtg atg ggg gac aaa tgt	1540
Ser Tyr Gly Arg Cys Ser Cys Lys Pro Gly Val Met Gly Asp Lys Cys	
415 420 425	
gac cgt tgc cag cct gga ttc cat tct ctc act gaa gca gga tgc agg	1588
Asp Arg Cys Gln Pro Gly Phe His Ser Leu Thr Glu Ala Gly Cys Arg	
430 435 440	
cca tgc tct tgt gat ccc tct ggc agc ata gat gaa tgt aat gtt gaa	1636
Pro Cys Ser Cys Asp Pro Ser Gly Ser Ile Asp Glu Cys Asn Val Glu	
445 450 455	
aca gga aga tgt gtt tgc aaa gac aat gtc gaa ggc ttc aat tgt gaa	1684
Thr Gly Arg Cys Val Cys Lys Asp Asn Val Glu Gly Phe Asn Cys Glu	
460 465 470 475	
aga tgc aaa cct gga ttt ttt aat ctg gaa tca tct aat cct cgg ggt	1732
Arg Cys Lys Pro Gly Phe Phe Asn Leu Glu Ser Ser Asn Pro Arg Gly	
480 485 490	
tgc aca ccc tgc ttc tgc ttt ggg cat tct tct gtc tgt aca aac gct	1780
Cys Thr Pro Cys Phe Cys Phe Gly His Ser Ser Val Cys Thr Asn Ala	
495 500 505	
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Val Gly Tyr Ser Val Tyr Ser Ile Ser Ser Thr Phe Gln Ile Asp Glu	
510 515 520	
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Asp Gly Trp Arg Ala Glu Gln Arg Asp Gly Ser Glu Ala Ser Leu Glu	
525 530 535	
tgg tcc tct gag agg caa gat atc gcc gtg atc tca gac agc tac ttt	1924
Trp Ser Ser Glu Arg Gln Asp Ile Ala Val Ile Ser Asp Ser Tyr Phe	
540 545 550 555	
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Pro Arg Tyr Phe Ile Ala Pro Ala Lys Phe Leu Gly Lys Gln Val Leu	
560 565 570	
agt tat ggt cag aac ctc tcc ttc tcc ttt cga gtg gac agg cga gat	2020
Ser Tyr Gly Gln Asn Leu Ser Phe Ser Phe Arg Val Asp Arg Arg Asp	
575 580 585	
act cgc ctc tct gcc gaa gac ctt gtg ctt gag gga gct ggc tta aga	2068
Thr Arg Leu Ser Ala Glu Asp Leu Val Leu Glu Gly Ala Gly Leu Arg	
590 595 600	
gta tct gta ccc ttg atc gct cag ggc aat tcc tat cca agt gag acc	2116
Val Ser Val Pro Leu Ile Ala Gln Gly Asn Ser Tyr Pro Ser Glu Thr	
605 610 615	
act gtg aag tat gtc ttc agg ctc cat gaa gca aca gat tac cct tgg	2164
Thr Val Lys Tyr Val Phe Arg Leu His Glu Ala Thr Asp Tyr Pro Trp	
620 625 630 635	

agg cct gct ctt acc cct ttt gaa ttt cag aag ctc cta aac aac ttg	2212
Arg Pro Ala Leu Thr Pro Phe Glu Phe Gln Lys Leu Leu Asn Asn Leu	
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acc tct atc aag ata cgt ggg aca tac agt gag aga agt gct gga tat	2260
Thr Ser Ile Lys Ile Arg Gly Thr Tyr Ser Glu Arg Ser Ala Gly Tyr	
655 660 665	
ttg gat gat gtc acc ctg gca agt gct cgt cct ggg cct gga gtc cct	2308
Leu Asp Asp Val Thr Leu Ala Ser Ala Arg Pro Gly Pro Gly Val Pro	
670 675 680	
gca act tgg gtg gag tcc tgc acc tgt cct gtg gga tat gga ggg cag	2356
Ala Thr Trp Val Glu Ser Cys Thr Cys Pro Val Gly Tyr Gly Gly Gln	
685 690 695	
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Phe Cys Glu Met Cys Leu Ser Gly Tyr Arg Arg Glu Thr Pro Asn Leu	
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gga cca tac agt cca tgt gtg ctt tgc gcc tgc aat gga cac agc gag	2452
Gly Pro Tyr Ser Pro Cys Val Leu Cys Ala Cys Asn Gly His Ser Glu	
720 725 730	
acc tgt gat cct gag aca ggt gtt tgt aac tgc aga gac aat acg gct	2500
Thr Cys Asp Pro Glu Thr Gly Val Cys Asn Cys Arg Asp Asn Thr Ala	
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ggc ccg cac tgt gag aag tgc agt gat ggg tac tat gga gat tca act	2548
Gly Pro His Cys Glu Lys Cys Ser Asp Gly Tyr Tyr Gly Asp Ser Thr	
750 755 760	
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Ala Gly Thr Ser Ser Asp Cys Gln Pro Cys Pro Cys Pro Gly Gly Ser	
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agt tgt gct gtt gtt ccc aag aca aag gag gtg gtg tgc acc aac tgt	2644
Ser Cys Ala Val Val Pro Lys Thr Lys Glu Val Val Cys Thr Asn Cys	
780 785 790 795	
cct act ggc acc act ggt aag aga tgt gag ctc tgt gat gat ggc tac	2692
Pro Thr Gly Thr Thr Gly Lys Arg Cys Glu Leu Cys Asp Asp Gly Tyr	
800 805 810	
ttt gga gac ccc ctg ggt aga aac ggc cct gtg aga ctt tgc cgc ctg	2740
Phe Gly Asp Pro Leu Gly Arg Asn Gly Pro Val Arg Leu Cys Arg Leu	
815 820 825	
tgc cag tgc agt gac aac atc gat ccc aac gca gtt gga aat tgc aat	2788
Cys Gln Cys Ser Asp Asn Ile Asp Pro Asn Ala Val Gly Asn Cys Asn	
830 835 840	
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Arg Leu Thr Gly Glu Cys Leu Lys Cys Ile Tyr Asn Thr Ala Gly Phe	
845 850 855	
tat tgt gac cgg tgc aaa gac gga ttt ttt gga aat ccc ctg gct ccc	2884
Tyr Cys Asp Arg Cys Lys Asp Gly Phe Phe Gly Asn Pro Leu Ala Pro	
860 865 870 875	
aat cca gca gac aaa tgc aaa gcc tgc aat tgc aat ccg tat ggg acc	2932

Asn	Pro	Ala	Asp	Lys	Cys	Lys	Ala	Cys	Asn	Cys	Asn	Pro	Tyr	Gly	Thr	
				880					885					890		
atg	aag	cag	cag	agc	agc	tgt	aac	ccc	gtg	acg	ggg	cag	tgt	gaa	tgt	2980
Met	Lys	Gln	Gln	Ser	Ser	Cys	Asn	Pro	Val	Thr	Gly	Gln	Cys	Glu	Cys	
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Leu	Pro	His	Val	Thr	Gly	Gln	Asp	Cys	Gly	Ala	Cys	Asp	Pro	Gly	Phe	
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Tyr	Asn	Leu	Gln	Ser	Gly	Gln	Gly	Cys	Glu	Arg	Cys	Asp	Cys	His	Ala	
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Leu	Gly	Ser	Thr	Asn	Gly	Gln	Cys	Asp	Ile	Arg	Thr	Gly	Gln	Cys	Glu	
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tgc	cag	ccc	ggc	atc	act	ggg	cag	cac	tgt	gag	cgc	tgt	gag	gtc	aac	3172
Cys	Gln	Pro	Gly	Ile	Thr	Gly	Gln	His	Cys	Glu	Arg	Cys	Glu	Val	Asn	
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cac	ttt	ggg	ttt	gga	cct	gaa	ggc	tgc	aaa	ccc	tgt	gac	tgt	cat	cct	3220
His	Phe	Gly	Phe	Gly	Pro	Glu	Gly	Cys	Lys	Pro	Cys	Asp	Cys	His	Pro	
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gag	gga	tct	ctt	tca	ctt	cag	tgc	aaa	gat	gat	ggg	cgc	tgt	gaa	tgc	3268
Glu	Gly	Ser	Leu	Ser	Leu	Gln	Cys	Lys	Asp	Asp	Gly	Arg	Cys	Glu	Cys	
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Arg	Glu	Gly	Phe	Val	Gly	Asn	Arg	Cys	Asp	Gln	Cys	Glu	Glu	Asn	Tyr	
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Phe	Tyr	Asn	Arg	Ser	Trp	Pro	Gly	Cys	Gln	Glu	Cys	Pro	Ala	Cys	Tyr	
1020					1025					1030				1035		
cgg	ctg	gta	aag	gat	aag	gtt	gct	gat	cat	aga	gtg	aag	ctc	cag	gaa	3412
Arg	Leu	Val	Lys	Asp	Lys	Val	Ala	Asp	His	Arg	Val	Lys	Leu	Gln	Glu	
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Leu	Glu	Ser	Leu	Ile	Ala	Asn	Leu	Gly	Thr	Gly	Asp	Glu	Met	Val	Thr	
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Asp	Gln	Ala	Phe	Glu	Asp	Arg	Leu	Lys	Glu	Ala	Glu	Arg	Glu	Val	Met	
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Asp	Leu	Leu	Arg	Glu	Ala	Gln	Asp	Val	Lys	Asp	Val	Asp	Gln	Asn	Leu	
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Met	Asp	Arg	Leu	Gln	Arg	Val	Asn	Asn	Thr	Leu	Ser	Ser	Gln	Ile	Ser	
1100					1105					1110				1115		
cgt	tta	cag	aat	atc	cgg	aat	acc	att	gaa	gag	act	gga	aac	ttg	gct	3652
Arg	Leu	Gln	Asn	Ile	Arg	Asn	Thr	Ile	Glu	Glu	Thr	Gly	Asn	Leu	Ala	

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	Glu Gln Ala Arg Ala His Val Glu Asn Thr Glu Arg Leu Ile Glu Ile			
	1135	1140	1145	
	gca tcc aga gaa ctt gag aaa gca aaa gtc gct gct gcc aat gtg tca			3748
	Ala Ser Arg Glu Leu Glu Lys Ala Lys Val Ala Ala Ala Asn Val Ser			
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	gtc act cag cca gaa tct aca ggg gac cca aac aac atg act ctt ttg			3796
	Val Thr Gln Pro Glu Ser Thr Gly Asp Pro Asn Asn Met Thr Leu Leu			
	1165	1170	1175	
	gca gaa gag gct cga aag ctt gct gaa cgt cat aaa cag gaa gct gat			3844
	Ala Glu Glu Ala Arg Lys Leu Ala Glu Arg His Lys Gln Glu Ala Asp			
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	gac att gtt cga gtg gca aag aca gcc aat gat acg tca act gag gca			3892
	Asp Ile Val Arg Val Ala Lys Thr Ala Asn Asp Thr Ser Thr Glu Ala			
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	Tyr Asn Leu Leu Leu Arg Thr Leu Ala Gly Glu Asn Gln Thr Ala Phe			
	1215	1220	1225	
	gag att gaa gag ctt aat agg aag tat gaa caa gcg aag aac atc tca			3988
	Glu Ile Glu Glu Leu Asn Arg Lys Tyr Glu Gln Ala Lys Asn Ile Ser			
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	cag gat ctg gaa aaa caa gct gcc cga gta cat gag gag gcc aaa agg			4036
	Gln Asp Leu Glu Lys Gln Ala Ala Arg Val His Glu Glu Ala Lys Arg			
	1245	1250	1255	
	gcc ggt gac aaa gct gtg gag atc tat gcc agc gtg gct cag ctg agc			4084
	Ala Gly Asp Lys Ala Val Glu Ile Tyr Ala Ser Val Ala Gln Leu Ser			
	1260	1265	1270	1275
	cct ttg gac tct gag aca ctg gag aat gaa gca aat aac ata aag atg			4132
	Pro Leu Asp Ser Glu Thr Leu Glu Asn Glu Ala Asn Asn Ile Lys Met			
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	gaa gct gag aat ctg gaa caa ctg att gac cag aaa tta aaa gat tat			4180
	Glu Ala Glu Asn Leu Glu Gln Leu Ile Asp Gln Lys Leu Lys Asp Tyr			
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	gag gac ctc aga gaa gat atg aga ggg aag gaa ctt gaa gtc aag aac			4228
	Glu Asp Leu Arg Glu Asp Met Arg Gly Lys Glu Leu Glu Val Lys Asn			
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	Leu Leu Glu Lys Gly Lys Thr Glu Gln Gln Thr Ala Asp Gln Leu Leu			
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Leu Gln Glu Ala Glu Lys Glu Leu Lys Arg Lys Gln Asp Asp Ala Asp	
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Gln Asp Met Met Met Ala Gly Met Ala Ser Gln Ala Ala Gln Glu Ala	
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Glu Ile Asn Ala Arg Lys Ala Lys Asn Ser Val Thr Ser Leu Leu Ser	
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 Arg Cys Met Pro Glu Phe Val Asn Ala Ala Phe Asn Val Thr Val Val
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 Ala Thr Asn Thr Cys Gly Thr Pro Pro Glu Glu Tyr Cys Val Gln Thr
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 Gly Val Thr Gly Val Thr Lys Ser Cys His Leu Cys Asp Ala Gly Gln
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 Pro His Leu Gln His Gly Ala Ala Phe Leu Thr Asp Tyr Asn Asn Gln
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 Ala Asp Thr Thr Trp Trp Gln Ser Gln Thr Met Leu Ala Gly Val Gln
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 Tyr Pro Ser Ser Ile Asn Leu Thr Leu His Leu Gly Lys Ala Phe Asp
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 Ile Thr Tyr Val Arg Leu Lys Phe His Thr Ser Arg Pro Glu Ser Phe
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 Ala Ile Tyr Lys Arg Thr Arg Glu Asp Gly Pro Trp Ile Pro Tyr Gln
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 Tyr Tyr Ser Gly Ser Cys Glu Asn Thr Tyr Ser Lys Ala Asn Arg Gly
 180 185 190
 Phe Ile Arg Thr Gly Gly Asp Glu Gln Gln Ala Leu Cys Thr Asp Glu
 195 200 205
 Phe Ser Asp Ile Ser Pro Leu Thr Gly Gly Asn Val Ala Phe Ser Thr
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 Leu Glu Gly Arg Pro Ser Ala Tyr Asn Phe Asp Asn Ser Pro Val Leu
 225 230 235 240
 Gln Glu Trp Val Thr Ala Thr Asp Ile Arg Val Thr Leu Asn Arg Leu

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Tyr	Tyr	Tyr	Ala	Ile	Ser	Asp	Phe	Ala	Val	Gly	Gly	Arg	Cys	Lys	Cys					
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Asn	Gly	His	Ala	Ser	Glu	Cys	Met	Lys	Asn	Glu	Phe	Asp	Lys	Leu	Val					
	290					295					300									
Cys	Asn	Cys	Lys	His	Asn	Thr	Tyr	Gly	Val	Asp	Cys	Glu	Lys	Cys	Leu					
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Pro	Phe	Phe	Asn	Asp	Arg	Pro	Trp	Arg	Arg	Ala	Thr	Ala	Glu	Ser	Ala					
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Ser	Glu	Cys	Leu	Pro	Cys	Asp	Cys	Asn	Gly	Arg	Ser	Gln	Glu	Cys	Tyr					
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Phe	Asp	Pro	Glu	Leu	Tyr	Arg	Ser	Thr	Gly	His	Gly	Gly	His	Cys	Thr					
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Asn	Cys	Gln	Asp	Asn	Thr	Asp	Gly	Ala	His	Cys	Glu	Arg	Cys	Arg	Glu					
	370					375					380									
Asn	Phe	Phe	Arg	Leu	Gly	Asn	Asn	Glu	Ala	Cys	Ser	Ser	Cys	His	Cys					
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Ser	Pro	Val	Gly	Ser	Leu	Ser	Thr	Gln	Cys	Asp	Ser	Tyr	Gly	Arg	Cys					
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Ser	Cys	Lys	Pro	Gly	Val	Met	Gly	Asp	Lys	Cys	Asp	Arg	Cys	Gln	Pro					
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Gly	Phe	His	Ser	Leu	Thr	Glu	Ala	Gly	Cys	Arg	Pro	Cys	Ser	Cys	Asp					
			435					440				445								
Pro	Ser	Gly	Ser	Ile	Asp	Glu	Cys	Asn	Val	Glu	Thr	Gly	Arg	Cys	Val					
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Cys	Phe	Gly	His	Ser	Ser	Val	Cys	Thr	Asn	Ala	Val	Gly	Tyr	Ser	Val					
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Glu	Gln	Arg	Asp	Gly	Ser	Glu	Ala	Ser	Leu	Glu	Trp	Ser	Ser	Glu	Arg					
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 Ile Ala Gln Gly Asn Ser Tyr Pro Ser Glu Thr Thr Val Lys Tyr Val
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 Phe Arg Leu His Glu Ala Thr Asp Tyr Pro Trp Arg Pro Ala Leu Thr
 625 630 635 640
 Pro Phe Glu Phe Gln Lys Leu Leu Asn Asn Leu Thr Ser Ile Lys Ile
 645 650 655
 Arg Gly Thr Tyr Ser Glu Arg Ser Ala Gly Tyr Leu Asp Asp Val Thr
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 Leu Ala Ser Ala Arg Pro Gly Pro Gly Val Pro Ala Thr Trp Val Glu
 675 680 685
 Ser Cys Thr Cys Pro Val Gly Tyr Gly Gly Gln Phe Cys Glu Met Cys
 690 695 700
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 705 710 715 720
 Cys Val Leu Cys Ala Cys Asn Gly His Ser Glu Thr Cys Asp Pro Glu
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 Thr Gly Val Cys Asn Cys Arg Asp Asn Thr Ala Gly Pro His Cys Glu
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 Lys Cys Ser Asp Gly Tyr Tyr Gly Asp Ser Thr Ala Gly Thr Ser Ser
 755 760 765
 Asp Cys Gln Pro Cys Pro Cys Pro Gly Gly Ser Ser Cys Ala Val Val
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 Pro Lys Thr Lys Glu Val Val Cys Thr Asn Cys Pro Thr Gly Thr Thr
 785 790 795 800
 Gly Lys Arg Cys Glu Leu Cys Asp Asp Gly Tyr Phe Gly Asp Pro Leu
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 Gly Arg Asn Gly Pro Val Arg Leu Cys Arg Leu Cys Gln Cys Ser Asp
 820 825 830
 Asn Ile Asp Pro Asn Ala Val Gly Asn Cys Asn Arg Leu Thr Gly Glu
 835 840 845
 Cys Leu Lys Cys Ile Tyr Asn Thr Ala Gly Phe Tyr Cys Asp Arg Cys
 850 855 860
 Lys Asp Gly Phe Phe Gly Asn Pro Leu Ala Pro Asn Pro Ala Asp Lys
 865 870 875 880
 Cys Lys Ala Cys Asn Cys Asn Pro Tyr Gly Thr Met Lys Gln Gln Ser
 885 890 895

Ser Cys Asn Pro Val Thr Gly Gln Cys Glu Cys Leu Pro His Val Thr
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Gly Gln Asp Cys Gly Ala Cys Asp Pro Gly Phe Tyr Asn Leu Gln Ser
 915 920 925

Gly Gln Gly Cys Glu Arg Cys Asp Cys His Ala Leu Gly Ser Thr Asn
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Gly Gln Cys Asp Ile Arg Thr Gly Gln Cys Glu Cys Gln Pro Gly Ile
 945 950 955 960

Thr Gly Gln His Cys Glu Arg Cys Glu Val Asn His Phe Gly Phe Gly
 965 970 975

Pro Glu Gly Cys Lys Pro Cys Asp Cys His Pro Glu Gly Ser Leu Ser
 980 985 990

Leu Gln Cys Lys Asp Asp Gly Arg Cys Glu Cys Arg Glu Gly Phe Val
 995 1000 1005

Gly Asn Arg Cys Asp Gln Cys Glu Glu Asn Tyr Phe Tyr Asn Arg Ser
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Trp Pro Gly Cys Gln Glu Cys Pro Ala Cys Tyr Arg Leu Val Lys Asp
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Lys Val Ala Asp His Arg Val Lys Leu Gln Glu Leu Glu Ser Leu Ile
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Ala Asn Leu Gly Thr Gly Asp Glu Met Val Thr Asp Gln Ala Phe Glu
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Asp Arg Leu Lys Glu Ala Glu Arg Glu Val Met Asp Leu Leu Arg Glu
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Ala Gln Asp Val Lys Asp Val Asp Gln Asn Leu Met Asp Arg Leu Gln
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Arg Val Asn Asn Thr Leu Ser Ser Gln Ile Ser Arg Leu Gln Asn Ile
 105 1110 1115 1120

Arg Asn Thr Ile Glu Glu Thr Gly Asn Leu Ala Glu Gln Ala Arg Ala
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His Val Glu Asn Thr Glu Arg Leu Ile Glu Ile Ala Ser Arg Glu Leu
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Glu Lys Ala Lys Val Ala Ala Ala Asn Val Ser Val Thr Gln Pro Glu
 1155 1160 1165

Ser Thr Gly Asp Pro Asn Asn Met Thr Leu Leu Ala Glu Glu Ala Arg
 1170 1175 1180

Lys Leu Ala Glu Arg His Lys Gln Glu Ala Asp Asp Ile Val Arg Val
 185 1190 1195 1200

Ala Lys Thr Ala Asn Asp Thr Ser Thr Glu Ala Tyr Asn Leu Leu Leu
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Arg Thr Leu Ala Gly Glu Asn Gln Thr Ala Phe Glu Ile Glu Glu Leu

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Ala Ser Ala Arg Pro Gly Pro Gly Val Pro Ala Thr Trp Val Glu Ser	
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Cys Thr Cys Pro Val Gly Tyr Gly Gly Gln Phe Cys Glu Met Cys Leu	
660 665 670	
tca ggt tac aga aga gaa act cct aat ctt gga cca tac agt cca tgt	2064
Ser Gly Tyr Arg Arg Glu Thr Pro Asn Leu Gly Pro Tyr Ser Pro Cys	
675 680 685	
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Val Leu Cys Ala Cys Asn Gly His Ser Glu Thr Cys Asp Pro Glu Thr	
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Gly Val Cys Asn Cys Arg Asp Asn Thr Ala Gly Pro His Cys Glu Lys	
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Cys Ser Asp Gly Tyr Tyr Gly Asp Ser Thr Ala Gly Thr Ser Ser Asp	
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Cys Gln Pro Cys Pro Cys Pro Gly Gly Ser Ser Cys Ala Val Val Pro	
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Lys Thr Lys Glu Val Val Cys Thr Asn Cys Pro Thr Gly Thr Thr Gly	
755 760 765	
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Lys Arg Cys Glu Leu Cys Asp Asp Gly Tyr Phe Gly Asp Pro Leu Gly	
770 775 780	
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Arg Asn Gly Pro Val Arg Leu Cys Arg Leu Cys Gln Cys Ser Asp Asn	
785 790 795 800	
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Ile Asp Pro Asn Ala Val Gly Asn Cys Asn Arg Leu Thr Gly Glu Cys	
805 810 815	
ctg aag tgc atc tat aac act gct ggc ttc tat tgt gac cgg tgc aaa	2496
Leu Lys Cys Ile Tyr Asn Thr Ala Gly Phe Tyr Cys Asp Arg Cys Lys	
820 825 830	
gac gga ttt ttt gga aat ccc ctg gct ccc aat cca gca gac aaa tgc	2544
Asp Gly Phe Phe Gly Asn Pro Leu Ala Pro Asn Pro Ala Asp Lys Cys	
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aaa gcc tgc aat tgc aat ccg tat ggg acc atg aag cag cag agc agc	2592
Lys Ala Cys Asn Cys Asn Pro Tyr Gly Thr Met Lys Gln Gln Ser Ser	
850 855 860	
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Cys Asn Pro Val Thr Gly Gln Cys Glu Cys Leu Pro His Val Thr Gly	
865 870 875 880	

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Gln Asp Cys Gly Ala Cys Asp Pro Gly Phe Tyr Asn Leu Gln Ser Gly	
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caa ggc tgt gag agg tgt gac tgc cat gcc ttg ggc tcc acc aat ggg	2736
Gln Gly Cys Glu Arg Cys Asp Cys His Ala Leu Gly Ser Thr Asn Gly	
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Gln Cys Asp Ile Arg Thr Gly Gln Cys Glu Cys Gln Pro Gly Ile Thr	
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Gly Gln His Cys Glu Arg Cys Glu Val Asn His Phe Gly Phe Gly Pro	
930 935 940	
gaa ggc tgc aaa ccc tgt gac tgt cat cct gag gga tct ctt tca ctt	2880
Glu Gly Cys Lys Pro Cys Asp Cys His Pro Glu Gly Ser Leu Ser Leu	
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Gln Cys Lys Asp Asp Gly Arg Cys Glu Cys Arg Glu Gly Phe Val Gly	
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Asn Arg Cys Asp Gln Cys Glu Glu Asn Tyr Phe Tyr Asn Arg Ser Trp	
980 985 990	
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Pro Gly Cys Gln Glu Cys Pro Ala Cys Tyr Arg Leu Val Lys Asp Lys	
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Val Ala Asp His Arg Val Lys Leu Gln Glu Leu Glu Ser Leu Ile Ala	
1010 1015 1020	
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Asn Leu Gly Thr Gly Asp Glu Met Val Thr Asp Gln Ala Phe Glu Asp	
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aga cta aag gaa gca gag agg gaa gtt atg gac ctc ctt cgt gag gcc	3168
Arg Leu Lys Glu Ala Glu Arg Glu Val Met Asp Leu Leu Arg Glu Ala	
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cag gat gtc aaa gat gtt gac cag aat ttg atg gat cgc cta cag aga	3216
Gln Asp Val Lys Asp Val Asp Gln Asn Leu Met Asp Arg Leu Gln Arg	
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Val Asn Asn Thr Leu Ser Ser Gln Ile Ser Arg Leu Gln Asn Ile Arg	
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Val Glu Asn Thr Glu Arg Leu Ile Glu Ile Ala Ser Arg Glu Leu Glu	
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Lys	Ala	Lys	Val	Ala	Ala	Ala	Asn	Val	Ser	Val	Thr	Gln	Pro	Glu	Ser		
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Thr	Gly	Asp	Pro	Asn	Asn	Met	Thr	Leu	Leu	Ala	Glu	Glu	Ala	Arg	Lys		
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Leu	Ala	Glu	Arg	His	Lys	Gln	Glu	Ala	Asp	Asp	Ile	Val	Arg	Val	Ala		
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Lys	Thr	Ala	Asn	Asp	Thr	Ser	Thr	Glu	Ala	Tyr	Asn	Leu	Leu	Leu	Arg		
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Arg	Lys	Tyr	Glu	Gln	Ala	Lys	Asn	Ile	Ser	Gln	Asp	Leu	Glu	Lys	Gln		
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Ala	Ala	Arg	Val	His	Glu	Glu	Ala	Lys	Arg	Ala	Gly	Asp	Lys	Ala	Val		
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gag	atc	tat	gcc	agc	gtg	gct	cag	ctg	agc	cct	ttg	gac	tct	gag	aca	3744	
Glu	Ile	Tyr	Ala	Ser	Val	Ala	Gln	Leu	Ser	Pro	Leu	Asp	Ser	Glu	Thr		
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ctg	gag	aat	gaa	gca	aat	aac	ata	aag	atg	gaa	gct	gag	aat	ctg	gaa	3792	
Leu	Glu	Asn	Glu	Ala	Asn	Asn	Ile	Lys	Met	Glu	Ala	Glu	Asn	Leu	Glu		
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caa	ctg	att	gac	cag	aaa	tta	aaa	gat	tat	gag	gac	ctc	aga	gaa	gat	3840	
Gln	Leu	Ile	Asp	Gln	Lys	Leu	Lys	Asp	Tyr	Glu	Asp	Leu	Arg	Glu	Asp		
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Met	Arg	Gly	Lys	Glu	Leu	Glu	Val	Lys	Asn	Leu	Leu	Glu	Lys	Gly	Lys		
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act	gaa	cag	cag	acc	gca	gac	caa	ctc	cta	gcc	cga	gct	gat	gct	gcc	3936	
Thr	Glu	Gln	Gln	Thr	Ala	Asp	Gln	Leu	Leu	Ala	Arg	Ala	Asp	Ala	Ala		
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Lys	Ala	Leu	Ala	Glu	Glu	Ala	Ala	Lys	Lys	Gly	Arg	Asp	Thr	Leu	Gln		
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gaa	gct	aat	gac	att	ctc	aac	aac	ctg	aaa	gat	ttt	gat	agg	cgc	gtg	4032	
Glu	Ala	Asn	Asp	Ile	Leu	Asn	Asn	Leu	Lys	Asp	Phe	Asp	Arg	Arg	Val		
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Ile	Asn	Gln	Thr	Ile	Thr	Glu	Ala	Asn	Glu	Lys	Thr	Arg	Glu	Ala	Gln		

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Gln Ala Leu Gly Ser Ala Ala Ala Asp Ala Thr Glu Ala Lys Asn Lys			
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gcc cat gag gcg gag agg atc gca agc gct gtc caa aag aat gcc acc			4224
Ala His Glu Ala Glu Arg Ile Ala Ser Ala Val Gln Lys Asn Ala Thr			
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agc acc aag gca gaa gct gaa aga act ttt gca gaa gtt aca gat ctg			4272
Ser Thr Lys Ala Glu Ala Glu Arg Thr Phe Ala Glu Val Thr Asp Leu			
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gat aat gag gtg aac aat atg ttg aag caa ctg cag gaa gca gaa aaa			4320
Asp Asn Glu Val Asn Asn Met Leu Lys Gln Leu Gln Glu Ala Glu Lys			
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Glu Leu Lys Arg Lys Gln Asp Asp Ala Asp Gln Asp Met Met Met Ala			
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ggg atg gct tca cag gct gct caa gaa gcc gag atc aat gcc aga aaa			4416
Gly Met Ala Ser Gln Ala Ala Gln Glu Ala Glu Ile Asn Ala Arg Lys			
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gcc aaa aac tct gtt act agc ctc ctc agc att att aat gac ctc ttg			4464
Ala Lys Asn Ser Val Thr Ser Leu Leu Ser Ile Ile Asn Asp Leu Leu			
1475	1480	1485	
gag cag ctg ggg cag ctg gat aca gtg gac ctg aat aag cta aac gag			4512
Glu Gln Leu Gly Gln Leu Asp Thr Val Asp Leu Asn Lys Leu Asn Glu			
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Ile Glu Gly Thr Leu Asn Lys Ala Lys Asp Glu Met Lys Val Ser Asp			
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ctt gat agg aaa gtg tct gac ctg gag aat gaa gcc aag aag cag gag			4608
Leu Asp Arg Lys Val Ser Asp Leu Glu Asn Glu Ala Lys Lys Gln Glu			
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gct gcc atc atg gac tat aac cga gat atc gag gag atc atg aag gac			4656
Ala Ala Ile Met Asp Tyr Asn Arg Asp Ile Glu Glu Ile Met Lys Asp			
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Ile Arg Asn Leu Glu Asp Ile Arg Lys Thr Leu Pro Ser Gly Cys Phe			
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Asn Thr Pro Ser Ile Glu Lys Pro			
1570	1575		
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<400> 24

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Cys	Met	Pro	Glu	Phe	Val	Asn	Ala	Ala	Phe	Asn	Val	Thr	Val	Val	Ala	20	25	30	
Thr	Asn	Thr	Cys	Gly	Thr	Pro	Pro	Glu	Glu	Tyr	Cys	Val	Gln	Thr	Gly	35	40	45	
Val	Thr	Gly	Val	Thr	Lys	Ser	Cys	His	Leu	Cys	Asp	Ala	Gly	Gln	Pro	50	55	60	
His	Leu	Gln	His	Gly	Ala	Ala	Phe	Leu	Thr	Asp	Tyr	Asn	Asn	Gln	Ala	65	70	75	80
Asp	Thr	Thr	Trp	Trp	Gln	Ser	Gln	Thr	Met	Leu	Ala	Gly	Val	Gln	Tyr	85	90	95	
Pro	Ser	Ser	Ile	Asn	Leu	Thr	Leu	His	Leu	Gly	Lys	Ala	Phe	Asp	Ile	100	105	110	
Thr	Tyr	Val	Arg	Leu	Lys	Phe	His	Thr	Ser	Arg	Pro	Glu	Ser	Phe	Ala	115	120	125	
Ile	Tyr	Lys	Arg	Thr	Arg	Glu	Asp	Gly	Pro	Trp	Ile	Pro	Tyr	Gln	Tyr	130	135	140	
Tyr	Ser	Gly	Ser	Cys	Glu	Asn	Thr	Tyr	Ser	Lys	Ala	Asn	Arg	Gly	Phe	145	150	155	160
Ile	Arg	Thr	Gly	Gly	Asp	Glu	Gln	Gln	Ala	Leu	Cys	Thr	Asp	Glu	Phe	165	170	175	
Ser	Asp	Ile	Ser	Pro	Leu	Thr	Gly	Gly	Asn	Val	Ala	Phe	Ser	Thr	Leu	180	185	190	
Glu	Gly	Arg	Pro	Ser	Ala	Tyr	Asn	Phe	Asp	Asn	Ser	Pro	Val	Leu	Gln	195	200	205	
Glu	Trp	Val	Thr	Ala	Thr	Asp	Ile	Arg	Val	Thr	Leu	Asn	Arg	Leu	Asn	210	215	220	
Thr	Phe	Gly	Asp	Glu	Val	Phe	Asn	Asp	Pro	Lys	Val	Leu	Lys	Ser	Tyr	225	230	235	240
Tyr	Tyr	Ala	Ile	Ser	Asp	Phe	Ala	Val	Gly	Gly	Arg	Cys	Lys	Cys	Asn	245	250	255	
Gly	His	Ala	Ser	Glu	Cys	Met	Lys	Asn	Glu	Phe	Asp	Lys	Leu	Val	Cys	260	265	270	
Asn	Cys	Lys	His	Asn	Thr	Tyr	Gly	Val	Asp	Cys	Glu	Lys	Cys	Leu	Pro	275	280	285	

Phe Phe Asn Asp Arg Pro Trp Arg Arg Ala Thr Ala Glu Ser Ala Ser
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 Glu Cys Leu Pro Cys Asp Cys Asn Gly Arg Ser Gln Glu Cys Tyr Phe
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 Cys Lys Pro Gly Val Met Gly Asp Lys Cys Asp Arg Cys Gln Pro Gly
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 Ser Gly Tyr Arg Arg Glu Thr Pro Asn Leu Gly Pro Tyr Ser Pro Cys
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 Gln Asp Cys Gly Ala Cys Asp Pro Gly Phe Tyr Asn Leu Gln Ser Gly
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 Gly Gln His Cys Glu Arg Cys Glu Val Asn His Phe Gly Phe Gly Pro

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Pro Gly Cys Gln Glu Cys Pro Ala Cys Tyr Arg Leu Val Lys Asp Lys		
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Asn Leu Gly Thr Gly Asp Glu Met Val Thr Asp Gln Ala Phe Glu Asp		
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Arg Leu Lys Glu Ala Glu Arg Glu Val Met Asp Leu Leu Arg Glu Ala		
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Gln Asp Val Lys Asp Val Asp Gln Asn Leu Met Asp Arg Leu Gln Arg		
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Asn Thr Ile Glu Glu Thr Gly Asn Leu Ala Glu Gln Ala Arg Ala His		
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Val Glu Asn Thr Glu Arg Leu Ile Glu Ile Ala Ser Arg Glu Leu Glu		
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Lys Ala Lys Val Ala Ala Ala Asn Val Ser Val Thr Gln Pro Glu Ser		
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Arg Lys Tyr Glu Gln Ala Lys Asn Ile Ser Gln Asp Leu Glu Lys Gln		
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Ala Ala Arg Val His Glu Glu Ala Lys Arg Ala Gly Asp Lys Ala Val		
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Glu Ile Tyr Ala Ser Val Ala Gln Leu Ser Pro Leu Asp Ser Glu Thr		
	1235	1240 1245
Leu Glu Asn Glu Ala Asn Asn Ile Lys Met Glu Ala Glu Asn Leu Glu		
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 1395 1400 1405
 Ser Thr Lys Ala Glu Ala Glu Arg Thr Phe Ala Glu Val Thr Asp Leu
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 Gly Met Ala Ser Gln Ala Ala Gln Glu Ala Glu Ile Asn Ala Arg Lys
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 cgccgcagcg gg atg acg ggc ggc ggg cgg gcc gcg ctg gcc ctg cag ccc 231
 Met Thr Gly Gly Gly Arg Ala Ala Leu Ala Leu Gln Pro
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 cgg ggg cgg ctg tgg ccg ctg ttg gct gtg ctg gcg gct gtg gcg ggc 279
 Arg Gly Arg Leu Trp Pro Leu Leu Ala Val Leu Ala Ala Val Ala Gly
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 Cys Val Arg Ala Ala Met Asp Glu Cys Ala Asp Glu Gly Gly Arg Pro
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 cag cgc tgc atg ccg gag ttt gtt aat gcc gcc ttc aat gtg acc gtg 375
 Gln Arg Cys Met Pro Glu Phe Val Asn Ala Ala Phe Asn Val Thr Val
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 gtg gct acc aac acg tgt ggg act ccg ccc gag gag tac tgc gtg cag 423
 Val Ala Thr Asn Thr Cys Gly Thr Pro Pro Glu Glu Tyr Cys Val Gln
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 act ggg gtg acc gga gtc act aag tcc tgt cac ctg tgc gac gcc ggc 471
 Thr Gly Val Thr Gly Val Thr Lys Ser Cys His Leu Cys Asp Ala Gly
 80 85 90
 cag cag cac ctg caa cac ggg gca gcc ttc ctg acc gac tac aac aac 519
 Gln Gln His Leu Gln His Gly Ala Ala Phe Leu Thr Asp Tyr Asn Asn
 95 100 105
 cag gcc gac acc acc tgg tgg caa agc cag act atg ctg gcc ggg gtg 567
 Gln Ala Asp Thr Thr Trp Trp Gln Ser Gln Thr Met Leu Ala Gly Val
 110 115 120 125
 cag tac ccc aac tcc atc aac ctc acg ctg cac ctg gga aag gct ttt 615
 Gln Tyr Pro Asn Ser Ile Asn Leu Thr Leu His Leu Gly Lys Ala Phe
 130 135 140
 gac atc act tac gtg cgc ctc aag ttc cac acc agc cgt cca gag agc 663
 Asp Ile Thr Tyr Val Arg Leu Lys Phe His Thr Ser Arg Pro Glu Ser
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Phe Ala Ile Tyr Lys Arg Thr Arg Glu Asp Gly Pro Trp Ile Pro Tyr	
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cag tac tac agt ggg tcc tgt gag aac acg tac tca aag gct aac cgt	759
Gln Tyr Tyr Ser Gly Ser Cys Glu Asn Thr Tyr Ser Lys Ala Asn Arg	
175 180 185	
ggc ttc atc agg acc gga ggg gac gag cag cag gcc ttg tgt act gat	807
Gly Phe Ile Arg Thr Gly Gly Asp Glu Gln Gln Ala Leu Cys Thr Asp	
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gaa ttc agt gac att tcc ccc ctc acc ggt ggc aac gtg gcc ttt tca	855
Glu Phe Ser Asp Ile Ser Pro Leu Thr Gly Gly Asn Val Ala Phe Ser	
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Thr Leu Glu Gly Arg Pro Ser Ala Tyr Asn Phe Asp Asn Ser Pro Val	
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Ser Tyr Tyr Tyr Ala Ile Ser Asp Phe Ala Val Gly Gly Arg Cys Lys	
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Cys Asn Gly His Ala Ser Glu Cys Val Lys Asn Glu Phe Asp Lys Leu	
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Met Cys Asn Cys Lys His Asn Thr Tyr Gly Val Asp Cys Glu Lys Cys	
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Leu Pro Phe Phe Asn Asp Arg Pro Trp Arg Arg Ala Thr Ala Glu Ser	
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Ala Ser Glu Cys Leu Pro Cys Asp Cys Asn Gly Arg Ser Gln Glu Cys	
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Tyr Phe Asp Pro Glu Leu Tyr Arg Ser Thr Gly His Gly Gly His Cys	
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Thr Asn Cys Arg Asp Asn Thr Asp Gly Ala Lys Cys Glu Arg Cys Arg	
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gag aat ttc ttc cgc ctg ggg aac act gaa gcc tgc tct ccg tgc cac	1383
Glu Asn Phe Phe Arg Leu Gly Asn Thr Glu Ala Cys Ser Pro Cys His	
385 390 395	
tgc agc cct gtt ggt tct ctc agc aca cag tgt gac agt tac ggc aga	1431

Cys	Ser	Pro	Val	Gly	Ser	Leu	Ser	Thr	Gln	Cys	Asp	Ser	Tyr	Gly	Arg	
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tgc	agc	tgt	aag	cca	gga	gtg	atg	ggc	gac	aag	tgt	gac	cgt	tgt	cag	1479
Cys	Ser	Cys	Lys	Pro	Gly	Val	Met	Gly	Asp	Lys	Cys	Asp	Arg	Cys	Gln	
		415				420					425					
cct	ggg	ttc	cat	tcc	ctc	act	gag	gca	gga	tgc	agg	cca	tgc	tcc	tgc	1527
Pro	Gly	Phe	His	Ser	Leu	Thr	Glu	Ala	Gly	Cys	Arg	Pro	Cys	Ser	Cys	
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gat	cct	tcg	ggc	agc	aca	gac	gag	tgt	aat	gtt	gaa	aca	gga	aga	tgc	1575
Asp	Pro	Ser	Gly	Ser	Thr	Asp	Glu	Cys	Asn	Val	Glu	Thr	Gly	Arg	Cys	
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gtt	tgc	aaa	gac	aat	gtt	gaa	ggc	ttc	aac	tgt	gag	aga	tgc	aaa	cct	1623
Val	Cys	Lys	Asp	Asn	Val	Glu	Gly	Phe	Asn	Cys	Glu	Arg	Cys	Lys	Pro	
			465					470					475			
gga	ttt	ttt	aat	ctg	gag	tca	tct	aat	cct	aag	ggc	tgc	aca	ccc	tgc	1671
Gly	Phe	Phe	Asn	Leu	Glu	Ser	Ser	Asn	Pro	Lys	Gly	Cys	Thr	Pro	Cys	
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ttc	tgc	ttt	ggc	cat	tct	tct	gtg	tgc	aca	aat	gct	gtt	ggc	tac	agt	1719
Phe	Cys	Phe	Gly	His	Ser	Ser	Val	Cys	Thr	Asn	Ala	Val	Gly	Tyr	Ser	
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gtt	tat	gac	atc	tcc	tcc	acc	ttt	cag	att	gat	gag	gat	ggg	tgg	cgc	1767
Val	Tyr	Asp	Ile	Ser	Ser	Thr	Phe	Gln	Ile	Asp	Glu	Asp	Gly	Trp	Arg	
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gtg	gag	cag	aga	gat	ggc	tcg	gag	gcg	tct	ctg	gag	tgg	tcc	tca	gac	1815
Val	Glu	Gln	Arg	Asp	Gly	Ser	Glu	Ala	Ser	Leu	Glu	Trp	Ser	Ser	Asp	
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agg	caa	tat	att	gcc	gta	atc	tca	gac	agt	tac	ttt	cct	aga	tac	ttc	1863
Arg	Gln	Tyr	Ile	Ala	Val	Ile	Ser	Asp	Ser	Tyr	Phe	Pro	Arg	Tyr	Phe	
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atc	gcc	cct	gtg	aag	ttc	ctg	ggc	aac	cag	gtc	ctg	agt	tat	ggg	cag	1911
Ile	Ala	Pro	Val	Lys	Phe	Leu	Gly	Asn	Gln	Val	Leu	Ser	Tyr	Gly	Gln	
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aat	ctt	tcc	ttc	tcc	ttc	cga	gtg	gac	aga	cga	gac	act	cgc	ctc	tcc	1959
Asn	Leu	Ser	Phe	Ser	Phe	Arg	Val	Asp	Arg	Arg	Asp	Thr	Arg	Leu	Ser	
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gca	gag	gac	ctt	gtg	ctc	gaa	gga	gct	ggc	ttg	aga	gta	tcc	gtg	ccc	2007
Ala	Glu	Asp	Leu	Val	Leu	Glu	Gly	Ala	Gly	Leu	Arg	Val	Ser	Val	Pro	
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Leu	Ile	Ala	Gln	Gly	Asn	Ser	Tyr	Pro	Ser	Glu	Thr	Thr	Val	Lys	Tyr	
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atc	ttc	agg	ctc	cat	gaa	gca	acg	gat	tac	cct	tgg	agg	ccc	gct	ctc	2103
Ile	Phe	Arg	Leu	His	Glu	Ala	Thr	Asp	Tyr	Pro	Trp	Arg	Pro	Ala	Leu	
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tcc	ccg	ttt	gaa	ttt	cag	aag	ctc	ctg	aac	aac	ttg	acc	tct	atc	aag	2151
Ser	Pro	Phe	Glu	Phe	Gln	Lys	Leu	Leu	Asn	Asn	Leu	Thr	Ser	Ile	Lys	

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ctg ggc agc aat ggg ccc gtg aga ctg tgc cgc ccg tgc cag tgt aac Leu Gly Ser Asn Gly Pro Val Arg Leu Cys Arg Pro Cys Gln Cys Asn 815 820 825			2679
gac aac ata gac ccc aac gcg gtt ggc aac tgc aac cgc ctg acg ggc Asp Asn Ile Asp Pro Asn Ala Val Gly Asn Cys Asn Arg Leu Thr Gly 830 835 840 845			2727
gag tgc ctg aag tgc atc tat aac acg gct ggt ttc tac tgc gac cgg Glu Cys Leu Lys Cys Ile Tyr Asn Thr Ala Gly Phe Tyr Cys Asp Arg 850 855 860			2775
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Ser	Cys	Asn	Pro	Val	Thr	Gly	Gln	Cys	Gln	Cys	Leu	Pro	His	Val	Ser	
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Gly	Arg	Asp	Cys	Gly	Thr	Cys	Asp	Pro	Gly	Tyr	Tyr	Asn	Leu	Gln	Ser	
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Thr	Gly	Gln	His	Cys	Glu	Arg	Cys	Glu	Thr	Asn	His	Phe	Gly	Phe	Gly	
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Pro	Glu	Gly	Cys	Lys	Pro	Cys	Asp	Cys	His	His	Glu	Gly	Ser	Leu	Ser	
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Leu	Gln	Cys	Lys	Asp	Asp	Gly	Arg	Cys	Glu	Cys	Arg	Glu	Gly	Phe	Val	
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Gly	Asn	Arg	Cys	Asp	Gln	Cys	Glu	Glu	Asn	Tyr	Phe	Tyr	Asn	Arg	Ser	
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Trp	Pro	Gly	Cys	Gln	Glu	Cys	Pro	Ala	Cys	Tyr	Arg	Leu	Val	Lys	Asp	
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aag	gct	gct	gag	cat	cga	gtg	aaa	ctc	cag	gag	tta	gag	agc	ctc	atc	3351
Lys	Ala	Ala	Glu	His	Arg	Val	Lys	Leu	Gln	Glu	Leu	Glu	Ser	Leu	Ile	
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Ala	Asn	Leu	Gly	Thr	Gly	Asp	Asp	Met	Val	Thr	Asp	Gln	Ala	Phe	Glu	
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Asp	Arg	Leu	Lys	Glu	Ala	Glu	Arg	Glu	Val	Thr	Asp	Leu	Leu	Arg	Glu	
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gct	cag	gaa	gtc	aaa	gat	gta	gat	caa	aat	ctg	atg	gat	cgc	ctt	cag	3495
Ala	Gln	Glu	Val	Lys	Asp	Val	Asp	Gln	Asn	Leu	Met	Asp	Arg	Leu	Gln	
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Lys Thr Ala Asn Glu Thr Ser Ala Glu Ala Tyr Asn Leu Leu Leu Arg	
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Arg Lys Tyr Glu Gln Ala Lys Asn Ile Ser Gln Asp Leu Glu Lys Gln	
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 gct gcc cga gtc cat gag gaa gcc aag cgt gca ggt gac aaa gcc gta	3975
Ala Ala Arg Val His Glu Glu Ala Lys Arg Ala Gly Asp Lys Ala Val	
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Glu Ile Tyr Ala Ser Val Ala Gln Leu Thr Pro Val Asp Ser Glu Ala	
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Arg Leu Ile Asp Gln Lys Leu Lys Asp Tyr Glu Asp Leu Arg Glu Asp	
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Met Arg Gly Lys Glu His Glu Val Lys Asn Leu Leu Glu Lys Gly Lys	
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Lys Ala Leu Ala Glu Glu Ala Ala Lys Lys Gly Arg Ser Thr Leu Gln	
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Glu Ala Asn Asp Ile Leu Asn Asn Leu Lys Asp Phe Asp Arg Arg Val	
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 Met Pro Glu Phe Val Asn Ala Ala Phe Asn Val Thr Val Val Ala Thr
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 Thr Gly Val Thr Lys Ser Cys His Leu Cys Asp Ala Gly Gln Gln His
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 Leu Gln His Gly Ala Ala Phe Leu Thr Asp Tyr Asn Asn Gln Ala Asp
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 Asn Ser Ile Asn Leu Thr Leu His Leu Gly Lys Ala Phe Asp Ile Thr
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 Tyr Val Arg Leu Lys Phe His Thr Ser Arg Pro Glu Ser Phe Ala Ile
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 Tyr Lys Arg Thr Arg Glu Asp Gly Pro Trp Ile Pro Tyr Gln Tyr Tyr

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Phe	Gly	Asp	Glu	Val	Phe	Asn	Asp	Pro	Lys	Val	Leu	Lys	Ser	Tyr	Tyr				
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 Gln Gly Asn Ser Tyr Pro Ser Glu Thr Thr Val Lys Tyr Ile Phe Arg
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 Gln Pro Cys Pro Cys Pro Gly Gly Ser Ser Cys Ala Ile Val Pro Lys
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 Lys Cys Ile Tyr Asn Thr Ala Gly Phe Tyr Cys Asp Arg Cys Lys Glu
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Arg His Lys Gln Glu Ala Asp Asp Ile Val Arg Val Ala Lys Thr Ala 1185	1190	1195 1200
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Lys Thr Ala Ala Glu Glu Ala Leu Arg Arg Ile Pro Ala Ile Asn Arg 1380	1385	1390
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 Pro Glu Phe Val Asn Ala Ala Phe Asn Val Thr Val Val Ala Thr Asn
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				165					170					175		
att	tcc	ccc	ctc	acc	ggg	ggc	aac	gtg	gcc	ttt	tca	acc	ctg	gaa	gga	576
Ile	Ser	Pro	Leu	Thr	Gly	Gly	Asn	Val	Ala	Phe	Ser	Thr	Leu	Glu	Gly	
			180					185					190			
cgg	ccg	agt	gcc	tac	aac	ttt	gac	aac	agc	cct	gtg	ctc	cag	gaa	tgg	624
Arg	Pro	Ser	Ala	Tyr	Asn	Phe	Asp	Asn	Ser	Pro	Val	Leu	Gln	Glu	Trp	
		195				200						205				
gta	act	gcc	act	gac	atc	aga	gtg	acg	ctc	aat	cgc	ctg	aac	acc	ttt	672
Val	Thr	Ala	Thr	Asp	Ile	Arg	Val	Thr	Leu	Asn	Arg	Leu	Asn	Thr	Phe	
	210					215					220					
gga	gat	gaa	gtg	ttt	aac	gac	ccc	aaa	gtt	ctc	aag	tct	tac	tat	tac	720
Gly	Asp	Glu	Val	Phe	Asn	Asp	Pro	Lys	Val	Leu	Lys	Ser	Tyr	Tyr	Tyr	
225					230					235					240	
gca	atc	tca	gac	ttt	gct	gtg	ggc	ggc	agg	tgt	aaa	tgt	aac	gga	cat	768
Ala	Ile	Ser	Asp	Phe	Ala	Val	Gly	Gly	Arg	Cys	Lys	Cys	Asn	Gly	His	
				245				250						255		
gcc	agc	gag	tgt	gta	aag	aac	gag	ttt	gac	aaa	ctc	atg	tgc	aac	tgc	816
Ala	Ser	Glu	Cys	Val	Lys	Asn	Glu	Phe	Asp	Lys	Leu	Met	Cys	Asn	Cys	
			260					265					270			
aaa	cat	aac	aca	tac	gga	gtt	gac	tgt	gaa	aag	tgc	ctg	cct	ttc	ttc	864
Lys	His	Asn	Thr	Tyr	Gly	Val	Asp	Cys	Glu	Lys	Cys	Leu	Pro	Phe	Phe	
		275					280					285				
aat	gac	cgg	ccg	tgg	agg	agg	gcg	act	gct	gag	agc	gcc	agc	gag	tgc	912
Asn	Asp	Arg	Pro	Trp	Arg	Arg	Ala	Thr	Ala	Glu	Ser	Ala	Ser	Glu	Cys	
	290					295					300					
ctt	cct	tgt	gac	tgc	aat	ggc	cga	tcc	caa	gag	tgc	tac	ttt	gat	cct	960
Leu	Pro	Cys	Asp	Cys	Asn	Gly	Arg	Ser	Gln	Glu	Cys	Tyr	Phe	Asp	Pro	
305					310					315					320	
gaa	cta	tac	cgt	tcc	act	gga	cat	ggg	ggc	cac	tgt	acc	aac	tgc	cgg	1008
Glu	Leu	Tyr	Arg	Ser	Thr	Gly	His	Gly	Gly	His	Cys	Thr	Asn	Cys	Arg	
				325				330						335		

gat aac aca gat ggt gcc aag tgc gag agg tgc cgg gag aat ttc ttc	1056
Asp Asn Thr Asp Gly Ala Lys Cys Glu Arg Cys Arg Glu Asn Phe Phe	
340 345 350	
cgc ctg ggg aac act gaa gcc tgc tct ccg tgc cac tgc agc cct gtt	1104
Arg Leu Gly Asn Thr Glu Ala Cys Ser Pro Cys His Cys Ser Pro Val	
355 360 365	
ggt tct ctc agc aca cag tgt gac agt tac ggc aga tgc agc tgt aag	1152
Gly Ser Leu Ser Thr Gln Cys Asp Ser Tyr Gly Arg Cys Ser Cys Lys	
370 375 380	
cca gga gtg atg ggt gac aag tgt gac cgt tgt cag cct ggg ttc cat	1200
Pro Gly Val Met Gly Asp Lys Cys Asp Arg Cys Gln Pro Gly Phe His	
385 390 395 400	
tcc ctc act gag gca gga tgc agg cca tgc tcc tgc gat cct tcg ggc	1248
Ser Leu Thr Glu Ala Gly Cys Arg Pro Cys Ser Cys Asp Pro Ser Gly	
405 410 415	
agc aca gac gag tgt aat gtt gaa aca gga aga tgc gtt tgc aaa gac	1296
Ser Thr Asp Glu Cys Asn Val Glu Thr Gly Arg Cys Val Cys Lys Asp	
420 425 430	
aat gtt gaa ggc ttc aac tgt gag aga tgc aaa cct gga ttt ttt aat	1344
Asn Val Glu Gly Phe Asn Cys Glu Arg Cys Lys Pro Gly Phe Phe Asn	
435 440 445	
ctg gag tca tct aat cct aag ggc tgc aca ccc tgc ttc tgc ttt ggc	1392
Leu Glu Ser Ser Asn Pro Lys Gly Cys Thr Pro Cys Phe Cys Phe Gly	
450 455 460	
cat tct tct gtg tgc aca aat gct gtt ggc tac agt gtt tat gac atc	1440
His Ser Ser Val Cys Thr Asn Ala Val Gly Tyr Ser Val Tyr Asp Ile	
465 470 475 480	
tcc tcc acc ttt cag att gat gag gat ggg tgg cgc gtg gag cag aga	1488
Ser Ser Thr Phe Gln Ile Asp Glu Asp Gly Trp Arg Val Glu Gln Arg	
485 490 495	
gat ggc tcg gag gcg tct ctg gag tgg tcc tca gac agg caa tat att	1536
Asp Gly Ser Glu Ala Ser Leu Glu Trp Ser Ser Asp Arg Gln Tyr Ile	
500 505 510	
gcc gta atc tca gac agt tac ttt cct aga tac ttc atc gcc cct gtg	1584
Ala Val Ile Ser Asp Ser Tyr Phe Pro Arg Tyr Phe Ile Ala Pro Val	
515 520 525	
aag ttc ctg ggc aac cag gtc ctg agt tat ggg cag aat ctt tcc ttc	1632
Lys Phe Leu Gly Asn Gln Val Leu Ser Tyr Gly Gln Asn Leu Ser Phe	
530 535 540	
tcc ttc cga gtg gac aga cga gac act cgc ctc tcc gca gag gac ctt	1680
Ser Phe Arg Val Asp Arg Arg Asp Thr Arg Leu Ser Ala Glu Asp Leu	
545 550 555 560	
gtg ctc gaa gga gct ggc ttg aga gta tcc gtg ccc ttg atc gct cag	1728
Val Leu Glu Gly Ala Gly Leu Arg Val Ser Val Pro Leu Ile Ala Gln	
565 570 575	

ggc aac tcc tac ccc agc gag acc act gtg aag tac atc ttc agg ctc	1776
Gly Asn Ser Tyr Pro Ser Glu Thr Thr Val Lys Tyr Ile Phe Arg Leu	
580 585 590	
cat gaa gca acg gat tac cct tgg agg ccc gct ctc tcc ccg ttt gaa	1824
His Glu Ala Thr Asp Tyr Pro Trp Arg Pro Ala Leu Ser Pro Phe Glu	
595 600 605	
ttt cag aag ctc ctg aac aac ttg acc tct atc aag atc cgt ggt aca	1872
Phe Gln Lys Leu Leu Asn Asn Leu Thr Ser Ile Lys Ile Arg Gly Thr	
610 615 620	
tac agc gag agg agc gct ggg tac ttg gat gat gtc acc ttg caa agt	1920
Tyr Ser Glu Arg Ser Ala Gly Tyr Leu Asp Asp Val Thr Leu Gln Ser	
625 630 635 640	
gct cgc cct ggg ccc gga gtc cct gca acg tgg gtg gag tcc tgc acc	1968
Ala Arg Pro Gly Pro Gly Val Pro Ala Thr Trp Val Glu Ser Cys Thr	
645 650 655	
tgt cca gtg gga tac ggg gga cag ttc tgt gag acg tgc ctc cca ggg	2016
Cys Pro Val Gly Tyr Gly Gly Gln Phe Cys Glu Thr Cys Leu Pro Gly	
660 665 670	
tac aga aga gaa act cca agc ctt gga cct tat agc ccg tgt gtg ctc	2064
Tyr Arg Arg Glu Thr Pro Ser Leu Gly Pro Tyr Ser Pro Cys Val Leu	
675 680 685	
tgt acc tgt aat ggg cac agt gag acc tgt gac ccg gag aca ggt gtc	2112
Cys Thr Cys Asn Gly His Ser Glu Thr Cys Asp Pro Glu Thr Gly Val	
690 695 700	
tgt gac tgc aga gac aat aca gcc ggc ccc cac tgt gag aaa tgt agc	2160
Cys Asp Cys Arg Asp Asn Thr Ala Gly Pro His Cys Glu Lys Cys Ser	
705 710 715 720	
gat ggg tac tat ggg gac tca acc ctg ggc acc tcc tct gac tgc cag	2208
Asp Gly Tyr Tyr Gly Asp Ser Thr Leu Gly Thr Ser Ser Asp Cys Gln	
725 730 735	
cct tgt ccc tgc ccc ggt ggc tca agt tgt gcc att gtc cca aag aca	2256
Pro Cys Pro Cys Pro Gly Gly Ser Ser Cys Ala Ile Val Pro Lys Thr	
740 745 750	
aag gaa gtg gtg tgc acg cac tgt ccg act ggc act gcc ggc aag aga	2304
Lys Glu Val Val Cys Thr His Cys Pro Thr Gly Thr Ala Gly Lys Arg	
755 760 765	
tgt gaa ctc tgt gat gac ggc tac ttt gga gac cct ctg ggc agc aat	2352
Cys Glu Leu Cys Asp Asp Gly Tyr Phe Gly Asp Pro Leu Gly Ser Asn	
770 775 780	
ggg ccc gtg aga ctg tgc cgc ccg tgc cag tgt aac gac aac ata gac	2400
Gly Pro Val Arg Leu Cys Arg Pro Cys Gln Cys Asn Asp Asn Ile Asp	
785 790 795 800	
ccc aac gcg gtt ggc aac tgc aac cgc ctg acg ggc gag tgc ctg aag	2448
Pro Asn Ala Val Gly Asn Cys Asn Arg Leu Thr Gly Glu Cys Leu Lys	
805 810 815	
tgc atc tat aac acg gct ggt ttc tac tgc gac cgg tgc aag gaa ggg	2496

Cys	Ile	Tyr	Asn	Thr	Ala	Gly	Phe	Tyr	Cys	Asp	Arg	Cys	Lys	Glu	Gly		
			820					825					830				
ttt	ttc	gga	aat	ccc	ctg	gct	ccc	aat	cca	gcc	gac	aaa	tgc	aaa	gcc	2544	
Phe	Phe	Gly	Asn	Pro	Leu	Ala	Pro	Asn	Pro	Ala	Asp	Lys	Cys	Lys	Ala		
		835					840					845					
tgc	gcc	tgc	aac	tac	ggg	aca	gtg	cag	caa	cag	agc	agc	tgt	aac	ccg	2592	
Cys	Ala	Cys	Asn	Tyr	Gly	Thr	Val	Gln	Gln	Gln	Ser	Ser	Cys	Asn	Pro		
	850					855					860						
gtg	acc	gga	caa	tgc	cag	tgt	ctg	cct	cat	gtg	tct	ggc	cgc	gac	tgc	2640	
Val	Thr	Gly	Gln	Cys	Gln	Cys	Leu	Pro	His	Val	Ser	Gly	Arg	Asp	Cys		
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ggt	act	tgt	gac	cct	ggc	tac	tac	aac	ctg	cag	agc	ggg	caa	ggc	tgc	2688	
Gly	Thr	Cys	Asp	Pro	Gly	Tyr	Tyr	Asn	Leu	Gln	Ser	Gly	Gln	Gly	Cys		
				885					890					895			
gag	agg	tgt	gac	tgc	cat	gct	ttg	ggt	tcc	acc	aat	ggg	cag	tgt	gac	2736	
Glu	Arg	Cys	Asp	Cys	His	Ala	Leu	Gly	Ser	Thr	Asn	Gly	Gln	Cys	Asp		
			900					905					910				
atc	cgc	acc	ggg	cag	tgt	gag	tgc	cag	cct	ggc	atc	acc	ggt	cag	cac	2784	
Ile	Arg	Thr	Gly	Gln	Cys	Glu	Cys	Gln	Pro	Gly	Ile	Thr	Gly	Gln	His		
		915					920					925					
tgt	gag	cgc	tgt	gag	acc	aac	cac	ttt	ggg	ttt	gga	cct	gaa	ggc	tgc	2832	
Cys	Glu	Arg	Cys	Glu	Thr	Asn	His	Phe	Gly	Phe	Gly	Pro	Glu	Gly	Cys		
	930					935					940						
aaa	cct	tgt	gac	tgt	cac	cat	gaa	gga	tcc	ctt	tcg	ctc	cag	tgt	aaa	2880	
Lys	Pro	Cys	Asp	Cys	His	His	Glu	Gly	Ser	Leu	Ser	Leu	Gln	Cys	Lys		
	945				950				955						960		
gac	gac	ggc	cgt	tgt	gaa	tgc	agg	gaa	ggc	ttt	gtg	ggc	aat	cgc	tgt	2928	
Asp	Asp	Gly	Arg	Cys	Glu	Cys	Arg	Glu	Gly	Phe	Val	Gly	Asn	Arg	Cys		
			965					970						975			
gac	cag	tgt	gaa	gag	aac	tat	ttc	tac	aat	cgg	tcc	tgg	cct	ggc	tgc	2976	
Asp	Gln	Cys	Glu	Glu	Asn	Tyr	Phe	Tyr	Asn	Arg	Ser	Trp	Pro	Gly	Cys		
		980						985					990				
cag	gag	tgt	ccg	gct	tgt	tac	cga	ctt	gtg	aag	gat	aag	gct	gct	gag	3024	
Gln	Glu	Cys	Pro	Ala	Cys	Tyr	Arg	Leu	Val	Lys	Asp	Lys	Ala	Ala	Glu		
		995				1000						1005					
cat	cga	gtg	aaa	ctc	cag	gag	tta	gag	agc	ctc	atc	gcc	aac	ctt	ggc	3072	
His	Arg	Val	Lys	Leu	Gln	Glu	Leu	Glu	Ser	Leu	Ile	Ala	Asn	Leu	Gly		
	1010				1015				1020								
act	ggg	gat	gac	atg	gtg	aca	gat	caa	gcc	ttt	gag	gac	aga	ctt	aag	3120	
Thr	Gly	Asp	Asp	Met	Val	Thr	Asp	Gln	Ala	Phe	Glu	Asp	Arg	Leu	Lys		
	1025				1030				1035					1040			
gaa	gca	gaa	agg	gag	gtg	aca	gac	ctt	ctc	cgt	gag	gct	cag	gaa	gtc	3168	
Glu	Ala	Glu	Arg	Glu	Val	Thr	Asp	Leu	Leu	Arg	Glu	Ala	Gln	Glu	Val		
			1045					1050					1055				
aaa	gat	gta	gat	caa	aat	ctg	atg	gat	cgc	ctt	cag	aga	gta	aat	agc	3216	
Lys	Asp	Val	Asp	Gln	Asn	Leu	Met	Asp	Arg	Leu	Gln	Arg	Val	Asn	Ser		

1060	1065	1070	
agc ctg cat agc caa att agc cga ctg cag aat atc cgg aat act atc			3264
Ser Leu His Ser Gln Ile Ser Arg Leu Gln Asn Ile Arg Asn Thr Ile			
1075	1080	1085	
gaa gag acc ggg atc ttg gct gag cga gca cgg tcc cga gtg gag agt			3312
Glu Glu Thr Gly Ile Leu Ala Glu Arg Ala Arg Ser Arg Val Glu Ser			
1090	1095	1100	
aca gag cag ctg att gag atc gcc tcc agg gag ctc gag aaa gca aaa			3360
Thr Glu Gln Leu Ile Glu Ile Ala Ser Arg Glu Leu Glu Lys Ala Lys			
1105	1110	1115	1120
atg gcc gcc aat gtg tca atc act cag cca gag tct aca ggg gag cca			3408
Met Ala Ala Asn Val Ser Ile Thr Gln Pro Glu Ser Thr Gly Glu Pro			
1125	1130	1135	
aac aac atg acc ctc ttg gca gaa gaa gcc cga agg ctt gca gag cgt			3456
Asn Asn Met Thr Leu Leu Ala Glu Glu Ala Arg Arg Leu Ala Glu Arg			
1140	1145	1150	
cat aaa cag gaa gcc gat gac att gta cga gtg gca aag aca gcc aac			3504
His Lys Gln Glu Ala Asp Asp Ile Val Arg Val Ala Lys Thr Ala Asn			
1155	1160	1165	
gag act tca gct gag gca tat aat ctg ctt ttg agg acc ctg gca gga			3552
Glu Thr Ser Ala Glu Ala Tyr Asn Leu Leu Leu Arg Thr Leu Ala Gly			
1170	1175	1180	
gaa aat caa act gcg ctg gag att gaa gaa ctt aac cgg aag tac gaa			3600
Glu Asn Gln Thr Ala Leu Glu Ile Glu Glu Leu Asn Arg Lys Tyr Glu			
1185	1190	1195	1200
caa gca aag aac atc tct cag gac ctg gag aag cag gct gcc cga gtc			3648
Gln Ala Lys Asn Ile Ser Gln Asp Leu Glu Lys Gln Ala Ala Arg Val			
1205	1210	1215	
cat gag gaa gcc aag cgt gca ggt gac aaa gcc gta gag atc tat gcc			3696
His Glu Glu Ala Lys Arg Ala Gly Asp Lys Ala Val Glu Ile Tyr Ala			
1220	1225	1230	
agt gtg gcc cag ctg acc cct gtg gac tct gag gcc ctg gag aat gaa			3744
Ser Val Ala Gln Leu Thr Pro Val Asp Ser Glu Ala Leu Glu Asn Glu			
1235	1240	1245	
gca aat aaa atc aag aaa gaa gct gca gac ctg gac cgt ctg att gac			3792
Ala Asn Lys Ile Lys Lys Glu Ala Ala Asp Leu Asp Arg Leu Ile Asp			
1250	1255	1260	
cag aag cta aag gat tac gag gac ctc agg gaa gac atg aga gga aag			3840
Gln Lys Leu Lys Asp Tyr Glu Asp Leu Arg Glu Asp Met Arg Gly Lys			
1265	1270	1275	1280
gaa cat gaa gtg aag aac ctt cta gag aag ggg aaa gct gaa cag cag			3888
Glu His Glu Val Lys Asn Leu Leu Glu Lys Gly Lys Ala Glu Gln Gln			
1285	1290	1295	
acc gcc gac caa ctc cta gct cga gcc gat gct gcc aag gcc ctt gct			3936
Thr Ala Asp Gln Leu Leu Ala Arg Ala Asp Ala Ala Lys Ala Leu Ala			
1300	1305	1310	

gaa gaa gct gct aag aag gga cgc agt acc tta caa gaa gcc aat gac	3984
Glu Glu Ala Ala Lys Lys Gly Arg Ser Thr Leu Gln Glu Ala Asn Asp	
1315 1320 1325	
att ctc aac aac ctg aaa gat ttt gat aga cgt gtg aac gat aac aag	4032
Ile Leu Asn Asn Leu Lys Asp Phe Asp Arg Arg Val Asn Asp Asn Lys	
1330 1335 1340	
aca gcc gcg gaa gaa gct cta agg aga att ccc gcc atc aac cgg acc	4080
Thr Ala Ala Glu Glu Ala Leu Arg Arg Ile Pro Ala Ile Asn Arg Thr	
1345 1350 1355 1360	
ata gct gaa gcc aat gag aag aca agg gag gcc cag cta gcg ctg ggc	4128
Ile Ala Glu Ala Asn Glu Lys Thr Arg Glu Ala Gln Leu Ala Leu Gly	
1365 1370 1375	
aat gct gcc gct gac gcc acg gag gcc aag aac aag gcc cat gag gca	4176
Asn Ala Ala Ala Asp Ala Thr Glu Ala Lys Asn Lys Ala His Glu Ala	
1380 1385 1390	
gag agg atc gcc agc gcc gcg cag aag aat gcc acc agt acc aag gcg	4224
Glu Arg Ile Ala Ser Ala Ala Gln Lys Asn Ala Thr Ser Thr Lys Ala	
1395 1400 1405	
gac gca gaa aga acc ttc ggg gaa gtt aca gat ctg gat aat gag gtg	4272
Asp Ala Glu Arg Thr Phe Gly Glu Val Thr Asp Leu Asp Asn Glu Val	
1410 1415 1420	
aac ggt atg ctg agg cag cta gag gag gca gag aat gag ctg aag agg	4320
Asn Gly Met Leu Arg Gln Leu Glu Glu Ala Glu Asn Glu Leu Lys Arg	
1425 1430 1435 1440	
aag caa gat gac gcc gac cag gac atg atg atg gcg ggg atg gct tcg	4368
Lys Gln Asp Asp Ala Asp Gln Asp Met Met Met Ala Gly Met Ala Ser	
1445 1450 1455	
caa gcc gct cag gag gct gag ctc aat gcc aga aag gcc aaa aac tct	4416
Gln Ala Ala Gln Glu Ala Glu Leu Asn Ala Arg Lys Ala Lys Asn Ser	
1460 1465 1470	
gtc agc agc ctc ctc agc cag ctg aac aac ctc ttg gat cag cta gga	4464
Val Ser Ser Leu Leu Ser Gln Leu Asn Asn Leu Leu Asp Gln Leu Gly	
1475 1480 1485	
cag ctg gac aca gtg gac ctg aac aag ctc aat gag atc gaa ggc tcc	4512
Gln Leu Asp Thr Val Asp Leu Asn Lys Leu Asn Glu Ile Glu Gly Ser	
1490 1495 1500	
ctg aac aaa gcc aaa gac gaa atg aag gcc agc gac ctg gac agg aag	4560
Leu Asn Lys Ala Lys Asp Glu Met Lys Ala Ser Asp Leu Asp Arg Lys	
1505 1510 1515 1520	
gtg tct gac ctg gag agc gag gct cgg aag cag gaa gca gcc atc atg	4608
Val Ser Asp Leu Glu Ser Glu Ala Arg Lys Gln Glu Ala Ala Ile Met	
1525 1530 1535	
gac tat aac cgg gac ata gca gag atc att aag gat att cac aac ctg	4656
Asp Tyr Asn Arg Asp Ile Ala Glu Ile Ile Lys Asp Ile His Asn Leu	
1540 1545 1550	

gag gac atc aag aag acc cta cca acc ggc tgc ttc aac acc ccg tct 4704
Glu Asp Ile Lys Lys Thr Leu Pro Thr Gly Cys Phe Asn Thr Pro Ser
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atc gag aag ccc tagtggcgag agggctgtaa ggcagtgtcc ctgacagggg 4756
Ile Glu Lys Pro
1570

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<212> PRT

<213> Mus musculus

<400> 28

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Pro	Glu	Phe	Val	Asn	Ala	Ala	Phe	Asn	Val	Thr	Val	Val	Ala	Thr	Asn
			20					25						30	
Thr	Cys	Gly	Thr	Pro	Pro	Glu	Glu	Tyr	Cys	Val	Gln	Thr	Gly	Val	Thr
		35					40					45			
Gly	Val	Thr	Lys	Ser	Cys	His	Leu	Cys	Asp	Ala	Gly	Gln	Gln	His	Leu
	50					55					60				
Gln	His	Gly	Ala	Ala	Phe	Leu	Thr	Asp	Tyr	Asn	Asn	Gln	Ala	Asp	Thr
	65				70					75				80	
Thr	Trp	Trp	Gln	Ser	Gln	Thr	Met	Leu	Ala	Gly	Val	Gln	Tyr	Pro	Asn
				85					90					95	
Ser	Ile	Asn	Leu	Thr	Leu	His	Leu	Gly	Lys	Ala	Phe	Asp	Ile	Thr	Tyr
			100					105					110		
Val	Arg	Leu	Lys	Phe	His	Thr	Ser	Arg	Pro	Glu	Ser	Phe	Ala	Ile	Tyr
			115				120					125			

Lys Arg Thr Arg Glu Asp Gly Pro Trp Ile Pro Tyr Gln Tyr Tyr Ser
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 Gly Ser Cys Glu Asn Thr Tyr Ser Lys Ala Asn Arg Gly Phe Ile Arg
 145 150 155 160
 Thr Gly Gly Asp Glu Gln Gln Ala Leu Cys Thr Asp Glu Phe Ser Asp
 165 170 175
 Ile Ser Pro Leu Thr Gly Gly Asn Val Ala Phe Ser Thr Leu Glu Gly
 180 185 190
 Arg Pro Ser Ala Tyr Asn Phe Asp Asn Ser Pro Val Leu Gln Glu Trp
 195 200 205
 Val Thr Ala Thr Asp Ile Arg Val Thr Leu Asn Arg Leu Asn Thr Phe
 210 215 220
 Gly Asp Glu Val Phe Asn Asp Pro Lys Val Leu Lys Ser Tyr Tyr Tyr
 225 230 235 240
 Ala Ile Ser Asp Phe Ala Val Gly Gly Arg Cys Lys Cys Asn Gly His
 245 250 255
 Ala Ser Glu Cys Val Lys Asn Glu Phe Asp Lys Leu Met Cys Asn Cys
 260 265 270
 Lys His Asn Thr Tyr Gly Val Asp Cys Glu Lys Cys Leu Pro Phe Phe
 275 280 285
 Asn Asp Arg Pro Trp Arg Arg Ala Thr Ala Glu Ser Ala Ser Glu Cys
 290 295 300
 Leu Pro Cys Asp Cys Asn Gly Arg Ser Gln Glu Cys Tyr Phe Asp Pro
 305 310 315 320
 Glu Leu Tyr Arg Ser Thr Gly His Gly Gly His Cys Thr Asn Cys Arg
 325 330 335
 Asp Asn Thr Asp Gly Ala Lys Cys Glu Arg Cys Arg Glu Asn Phe Phe
 340 345 350
 Arg Leu Gly Asn Thr Glu Ala Cys Ser Pro Cys His Cys Ser Pro Val
 355 360 365
 Gly Ser Leu Ser Thr Gln Cys Asp Ser Tyr Gly Arg Cys Ser Cys Lys
 370 375 380
 Pro Gly Val Met Gly Asp Lys Cys Asp Arg Cys Gln Pro Gly Phe His
 385 390 395 400
 Ser Leu Thr Glu Ala Gly Cys Arg Pro Cys Ser Cys Asp Pro Ser Gly
 405 410 415
 Ser Thr Asp Glu Cys Asn Val Glu Thr Gly Arg Cys Val Cys Lys Asp
 420 425 430
 Asn Val Glu Gly Phe Asn Cys Glu Arg Cys Lys Pro Gly Phe Phe Asn
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Leu Glu Ser Ser Asn Pro Lys Gly Cys Thr Pro Cys Phe Cys Phe Gly
 450 455 460
 His Ser Ser Val Cys Thr Asn Ala Val Gly Tyr Ser Val Tyr Asp Ile
 465 470 475 480
 Ser Ser Thr Phe Gln Ile Asp Glu Asp Gly Trp Arg Val Glu Gln Arg
 485 490 495
 Asp Gly Ser Glu Ala Ser Leu Glu Trp Ser Ser Asp Arg Gln Tyr Ile
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 Ala Val Ile Ser Asp Ser Tyr Phe Pro Arg Tyr Phe Ile Ala Pro Val
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 Ser Phe Arg Val Asp Arg Arg Asp Thr Arg Leu Ser Ala Glu Asp Leu
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 Val Leu Glu Gly Ala Gly Leu Arg Val Ser Val Pro Leu Ile Ala Gln
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 Gly Asn Ser Tyr Pro Ser Glu Thr Thr Val Lys Tyr Ile Phe Arg Leu
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 His Glu Ala Thr Asp Tyr Pro Trp Arg Pro Ala Leu Ser Pro Phe Glu
 595 600 605
 Phe Gln Lys Leu Leu Asn Asn Leu Thr Ser Ile Lys Ile Arg Gly Thr
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 Tyr Ser Glu Arg Ser Ala Gly Tyr Leu Asp Asp Val Thr Leu Gln Ser
 625 630 635 640
 Ala Arg Pro Gly Pro Gly Val Pro Ala Thr Trp Val Glu Ser Cys Thr
 645 650 655
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 Tyr Arg Arg Glu Thr Pro Ser Leu Gly Pro Tyr Ser Pro Cys Val Leu
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 690 695 700
 Cys Asp Cys Arg Asp Asn Thr Ala Gly Pro His Cys Glu Lys Cys Ser
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 Asp Gly Tyr Tyr Gly Asp Ser Thr Leu Gly Thr Ser Ser Asp Cys Gln
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 Pro Cys Pro Cys Pro Gly Gly Ser Ser Cys Ala Ile Val Pro Lys Thr
 740 745 750
 Lys Glu Val Val Cys Thr His Cys Pro Thr Gly Thr Ala Gly Lys Arg
 755 760 765
 Cys Glu Leu Cys Asp Asp Gly Tyr Phe Gly Asp Pro Leu Gly Ser Asn

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Pro Asn Ala Val Gly Asn Cys Asn Arg Leu Thr Gly Glu Cys Leu Lys 805 810 815		
Cys Ile Tyr Asn Thr Ala Gly Phe Tyr Cys Asp Arg Cys Lys Glu Gly 820 825 830		
Phe Phe Gly Asn Pro Leu Ala Pro Asn Pro Ala Asp Lys Cys Lys Ala 835 840 845		
Cys Ala Cys Asn Tyr Gly Thr Val Gln Gln Gln Ser Ser Cys Asn Pro 850 855 860		
Val Thr Gly Gln Cys Gln Cys Leu Pro His Val Ser Gly Arg Asp Cys 865 870 875 880		
Gly Thr Cys Asp Pro Gly Tyr Tyr Asn Leu Gln Ser Gly Gln Gly Cys 885 890 895		
Glu Arg Cys Asp Cys His Ala Leu Gly Ser Thr Asn Gly Gln Cys Asp 900 905 910		
Ile Arg Thr Gly Gln Cys Glu Cys Gln Pro Gly Ile Thr Gly Gln His 915 920 925		
Cys Glu Arg Cys Glu Thr Asn His Phe Gly Phe Gly Pro Glu Gly Cys 930 935 940		
Lys Pro Cys Asp Cys His His Glu Gly Ser Leu Ser Leu Gln Cys Lys 945 950 955 960		
Asp Asp Gly Arg Cys Glu Cys Arg Glu Gly Phe Val Gly Asn Arg Cys 965 970 975		
Asp Gln Cys Glu Glu Asn Tyr Phe Tyr Asn Arg Ser Trp Pro Gly Cys 980 985 990		
Gln Glu Cys Pro Ala Cys Tyr Arg Leu Val Lys Asp Lys Ala Ala Glu 995 1000 1005		
His Arg Val Lys Leu Gln Glu Leu Glu Ser Leu Ile Ala Asn Leu Gly 1010 1015 1020		
Thr Gly Asp Asp Met Val Thr Asp Gln Ala Phe Glu Asp Arg Leu Lys 1025 1030 1035 1040		
Glu Ala Glu Arg Glu Val Thr Asp Leu Leu Arg Glu Ala Gln Glu Val 1045 1050 1055		
Lys Asp Val Asp Gln Asn Leu Met Asp Arg Leu Gln Arg Val Asn Ser 1060 1065 1070		
Ser Leu His Ser Gln Ile Ser Arg Leu Gln Asn Ile Arg Asn Thr Ile 1075 1080 1085		
Glu Glu Thr Gly Ile Leu Ala Glu Arg Ala Arg Ser Arg Val Glu Ser 1090 1095 1100		

Thr Glu Gln Leu Ile Glu Ile Ala Ser Arg Glu Leu Glu Lys Ala Lys
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 Met Ala Ala Asn Val Ser Ile Thr Gln Pro Glu Ser Thr Gly Glu Pro
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 His Lys Gln Glu Ala Asp Asp Ile Val Arg Val Ala Lys Thr Ala Asn
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 Ser Val Ala Gln Leu Thr Pro Val Asp Ser Glu Ala Leu Glu Asn Glu
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 Glu His Glu Val Lys Asn Leu Leu Glu Lys Gly Lys Ala Glu Gln Gln
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 Thr Ala Asp Gln Leu Leu Ala Arg Ala Asp Ala Ala Lys Ala Leu Ala
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 Thr Ala Ala Glu Glu Ala Leu Arg Arg Ile Pro Ala Ile Asn Arg Thr
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 Ile Ala Glu Ala Asn Glu Lys Thr Arg Glu Ala Gln Leu Ala Leu Gly
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 Glu Arg Ile Ala Ser Ala Ala Gln Lys Asn Ala Thr Ser Thr Lys Ala
 1395 1400 1405
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Asn Gly Met Leu Arg Gln Leu Glu Glu Ala Glu Asn Glu Leu Lys Arg
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Lys Gln Asp Asp Ala Asp Gln Asp Met Met Met Ala Gly Met Ala Ser
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Gln Ala Ala Gln Glu Ala Glu Leu Asn Ala Arg Lys Ala Lys Asn Ser
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Val Ser Ser Leu Leu Ser Gln Leu Asn Asn Leu Leu Asp Gln Leu Gly
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Asp Tyr Asn Arg Asp Ile Ala Glu Ile Ile Lys Asp Ile His Asn Leu
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Glu Asp Ile Lys Lys Thr Leu Pro Thr Gly Cys Phe Asn Thr Pro Ser
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Ile Glu Lys Pro
1570

PATENT COOPERATION TREATY

DEC 28 2000

From the INTERNATIONAL SEARCHING AUTHORITY

PCT

NOTIFICATION OF TRANSMITTAL OF
THE INTERNATIONAL SEARCH REPORT
OR THE DECLARATION

(PCT Rule 44.1)

To:

MCDONNELL BOEHNEN HULBERT
& BERGHOFF
Attn. HARPER, David S.
300 South Wacker Drive
Suite 3200
Chicago, IL 60606
UNITED STATES OF AMERICA

Date of mailing
(day/month/year)

27/12/2000

Applicant's or agent's file reference

99,274-D1

FOR FURTHER ACTION

See paragraphs 1 and 4 below

International application No.

PCT/US 00/11543

International filing date
(day/month/year)

28/04/2000

Applicant

BIOSTRATUM, INC. et al.

1. ☒ The applicant is hereby notified that the International Search Report has been established and is transmitted herewith.

Filing of amendments and statement under Article 19:

The applicant is entitled, if he so wishes, to amend the claims of the International Application (see Rule 46):

When? The time limit for filing such amendments is normally 2 months from the date of transmittal of the International Search Report; however, for more details, see the notes on the accompanying sheet.

Where? Directly to the International Bureau of WIPO
34, chemin des Colombettes
1211 Geneva 20, Switzerland
Facsimile No.: (41-22) 740.14.35

For more detailed instructions, see the notes on the accompanying sheet.

2. ☐ The applicant is hereby notified that no International Search Report will be established and that the declaration under Article 17(2)(a) to that effect is transmitted herewith.

3. ☐ With regard to the protest against payment of (an) additional fee(s) under Rule 40.2, the applicant is notified that:

☐ the protest together with the decision thereon has been transmitted to the International Bureau together with the applicant's request to forward the texts of both the protest and the decision thereon to the designated Offices.

☐ no decision has been made yet on the protest; the applicant will be notified as soon as a decision is made.

4. **Further action(s):** The applicant is reminded of the following:

Shortly after **18 months** from the priority date, the international application will be published by the International Bureau. If the applicant wishes to avoid or postpone publication, a notice of withdrawal of the international application, or of the priority claim, must reach the International Bureau as provided in Rules 90bis.1 and 90bis.3, respectively, before the completion of the technical preparations for international publication.

Within **19 months** from the priority date, a demand for international preliminary examination must be filed if the applicant wishes to postpone the entry into the national phase until 30 months from the priority date (in some Offices even later).

Within **20 months** from the priority date, the applicant must perform the prescribed acts for entry into the national phase before all designated Offices which have not been elected in the demand or in a later election within 19 months from the priority date or could not be elected because they are not bound by Chapter II.

Name and mailing address of the International Searching Authority



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Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Mireille Claudepierre